



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 151522

TO: Elizabeth McElwain
Location: REM-2A11/2C18
Art Unit: 1638
Thursday, April 28, 2005

Case Serial Number: 10/088079

From: Edward Hart
Location: Biotech-Chem Library
REM-1A55
Phone: 571-272-2512

edward.hart@uspto.gov

Search Notes

Examiner McElwain,

Here are the results of the search you requested.

Please feel free to contact me if you have any questions.

Edward Hart



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(jtdpsn) Kuabla ebad pad
This

STIC-Biotech/ChemLib

151522

From: McElwain, Elizabeth
Sent: Friday, April 22, 2005 3:50 PM
To: STIC-Biotech/ChemLib
Subject: sequence search

Please search 10/088,079 - SEQ ID NO: 1 and 2, and DNA encoding SEQ ID NO: 2
for prior art and for interference.

Thank you,
Beth

Elizabeth F. McElwain, Ph.D.
U.S. Patent and Trademark Office
Tech Center 1600, Art Unit 1638
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mailbox Remsen 2C18
571-272-0802
elizabeth.mcelwain@uspto.gov

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4/27/05
1- AA - 01
1- AA - 020
reverts to AA
4/28/05

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: April 27, 2005, 10:53:47 ; Search time 40 Seconds
(without alignments)
815.437 Million cell updates/sec

Title: US-10-088-079-2
Perfect score: 1722
Sequence: 1 MNQRNASMTVIGAGSYGTAL.....AREAAALTLGRARKDERSH 339

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues
Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 20000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 79: *
1: pir1: *
2: pir2: *
3: pir3: *
4: pir4: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1719	99.8	339	2	S47829 glycerol-3-phospha
2	1719	99.8	339	2	G86036 glycerol-3-phospha
3	1719	99.8	339	2	F91189 glycerol-3-phospha
4	1640	95.2	339	2	AB0975 glycerol-3-phospha
5	1455	84.5	339	2	AD0009 glycerol-3-phospha
6	1242	72.1	344	2	A82050 glycerol-3-phospha
7	1150	66.8	335	2	F64080 glycerol-3-phospha
8	751.5	43.6	346	2	H82637 glycerol-3-phospha
9	703	40.8	332	2	C97111 glycerol-3-phospha
10	672.5	39.1	338	2	AH1316 NAD(P)H-dependent
11	669.5	38.9	338	2	AH1688 NAD(P)H-dependent
12	666.5	38.7	345	2	H69636 glycerol-3-phospha
13	631	36.6	345	2	H83854 NAD(P)H-dependent
14	624.5	36.3	329	2	E81953 glycerol-3-phospha
15	621	36.1	336	2	T35643 glycerol-3-phospha
16	613.5	35.6	341	2	B86792 hypothetical prote
17	608.5	35.3	334	2	C70673 probable gpda2 pro
18	607	35.2	327	2	AI2901 glycerol-3-phospha
19	607	35.2	327	2	C97677 probable glycerol-
20	605.5	35.2	338	2	C98109 glycerol-3-phospha
21	604.5	35.1	338	2	E95244 glycerol-3-phospha
22	601.5	34.9	332	2	AI3273 glycerol-3-phospha
23	599.5	34.8	326	2	B89926 glycerol-3-phospha
24	575.5	33.4	331	2	E87257 glycerol-3-phospha
25	567	32.9	349	2	T45431 glycerol-3-phospha
26	567	32.9	351	2	A87119 glycerol-3-phospha
27	556	32.3	340	2	H83443 glycerol-3-phospha
28	547	31.8	334	2	A81743 glycerol-3-phospha
29	546	31.7	334	2	G72024 glycerol-3-phospha

30	546	31.7	334	2	E86597 glycerol-3-P dehyd
31	532	30.9	334	2	A71480 probable glycerol-
32	515	29.9	341	2	C70932 probable dehydroge
33	505	29.3	328	2	H75251 glycerol-3-phospha
34	502.5	29.2	313	2	A70441 glycerol-3-phospha
35	482	28.0	330	2	S75139 glycerol-3-phospha
36	460	26.7	354	2	T48649 glycerol-3-phospha
37	448	26.0	307	2	AG2017 glycerol-3-phospha
38	422.5	24.5	433	2	F84832 glycerol-3-phospha
39	415	24.1	356	2	E71252 probable glycerol-
40	403	23.4	312	2	H71876 glycerol-3-phospha
41	403	23.4	312	2	A64640 glycerol-3-phospha
42	381	22.1	298	2	F81325 glycerol-3-phospha
43	366.5	21.3	316	2	A71703 glycerol-3-phospha
44	350	20.3	321	2	E69147 glycerol-3-phospha
45	350	20.3	325	2	G97776 hypothetical prote

ALIGNMENTS

RESULT 1

S47829
glycerol-3-phosphate dehydrogenase (NAD) (EC 1.1.1.8) - Escherichia coli (strain K-12)
C;Species: Escherichia coli
C;Date: 27-Jan-1995 #sequence_revision 27-Jan-1995 #text_change 09-Jul-2004
C;Accession: S47829; B65161
R;Plunkett, G.
submitted to the EMBL Data Library, March 1994
A;Reference number: S47666
A;Accession: S47829
A;Molecule type: DNA
A;Residues: 1-339 <PLU>
A;Cross-references: UNIPROT:P37606; EMBL:U00039; NID:G466582; PIDN:AAB18585.1; PID:G1657
R;Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.; Co
.A.; Rose, D.J.; Mau, B.; Shao, Y.
Science 277, 1453-1462, 1997
A;Title: The complete genome sequence of Escherichia coli K-12.
A;Reference number: A64720; MUID:97426617; PMID:9278503
A;Accession: B65161
A;Status: preliminary; nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-339 <BLAT>
A;Cross-references: GB:AE000439; GB:U00096; NID:G1790036; PIDN:AAC76632.1; PID:G1790037;
A;Experimental source: strain K-12, substrain MG1655
C;Genetics:
A;Gene: gpsA
C;Superfamily: glycerol-3-phosphate dehydrogenase (NAD)
C;Keywords: oxidoreductase

Query Match 99.8%; Score 1719; DB 2; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.2e-118;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy	1	MNQRNASMTVIGAGSYGTALAITLARNGHEVVWGHDP	PEHIATLERDRCNAAFLPDVPPF	60
Db	1	MNQRNASMTVIGAGSYGTALAITLARNGHEVVWGHDP	PEHIATLERDRCNAAFLPDVPPF	60
Qy	61	DTLHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMR	PDARLVWATKGLEAETGRLL	120
Db	61	DTLHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMR	PDARLVWATKGLEAETGRLL	120
Qy	121	QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLAST	DTQTFADDLQQLHCGKSFVY	180
Db	121	QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLAST	DTQTFADDLQQLHCGKSFVY	180
Qy	181	SNPDFIGVQLGGAVKNVIAIGAGMSDGI	GFGANARTALITRGLAEMSR	LGALGADPATF 240
Db	181	SNPDFIGVQLGGAVKNVIAIGAGMSDGI	GFGANARTALITRGLAEMSR	LGALGADPATF 240
Qy	241	MGMAGLGLDLVLTCTENQSRNRRFGMMLGQGM	DVQSAQEKIGQVVEGYRNTKEVRELAH	RF 300
Db	241	MGMAGLGLDLVLTCTENQSRNRRFGMMLGQGM	DVQSAQEKIGQVVEGYRNTKEVRELAH	RF 300

Qy 301 GVEMPTIEIYQVLYCGKNAREAAALTLGLRARKDERSH 339
Db 301 GVEMPTIEIYQVLYCGKNAREAAALTLGLRARKDERSH 339

RESULT 2
G86036
glycerol-3-phosphate dehydrogenase (NAD+) [imported] - Escherichia coli (strain O157:H7,
C;Species: Escherichia coli
C;Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 09-Jul-2004
C;Accession: G86036
R;Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew
iller, L.; Grotbeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca,
Nature 409, 529-533, 2001
A;Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
A;Reference number: A85480; MUID:21074935; PMID:11206551
A;Accession: G86036
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-339 <STO>
A;Cross-references: UNIPROT:P37606; GB:AE005174; NID:G12518358; PIDN:AAG58755.1; GSPDB:G
A;Experimental source: strain O157:H7, substrain EDL933
C;Genetics:
A;Gene: gpsA
C;Superfamily: glycerol-3-phosphate dehydrogenase (NAD)

Query Match 99.8%; Score 1719; DB 2; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.2e-118;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60
Db 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60

Qy 61 DTLHLESDLATALAASRNILVVPSPHVFGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
Db 61 DTLHLESDLATALAASRNILVVPSPHVFGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120

Qy 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFRVY 180
Db 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFRVY 180

Qy 181 SNPDFIGVLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRIGALGADPATF 240
Db 181 SNPDFIGVLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRIGALGADPATF 240

Qy 241 MGMAGLDLVLTCCTENQSRNRRFGMLQGQMDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300
Db 241 MGMAGLDLVLTCCTENQSRNRRFGMLQGQMDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300

Qy 301 GVEMPTIEIYQVLYCGKNAREAAALTLGLRARKDERSH 339
Db 301 GVEMPTIEIYQVLYCGKNAREAAALTLGLRARKDERSH 339

RESULT 3
F91189
glycerol-3-phosphate dehydrogenase (NAD+) [imported] - Escherichia coli (strain O157:H7,
C;Species: Escherichia coli
C;Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 16-Aug-2004
C;Accession: F91189
R;Hayashi, T.; Makino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.G.
gasawara, N.; Yasunaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.
DNA Res. 8, 11-22, 2001
A;Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and genc
A;Reference number: A99629; MUID:21156231; PMID:11258796
A;Accession: F91189
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-339 <HAY>
A;Cross-references: UNIPROT:P37606; GB:BA000007; PIDN:BAB37909.1; PID:G13363961; GSPDB:G
A;Experimental source: strain O157:H7, substrain RIMD 0509952

C;Genetics:
A;Gene: ECs4486
C;Superfamily: Glycerol-3-phosphate dehydrogenase (NAD)

Query Match 99.8%; Score 1719; DB 2; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.2e-118;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60
Db 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60

Qy 61 DTLHLESDLATALAASRNILVVPSPHVFGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
Db 61 DTLHLESDLATALAASRNILVVPSPHVFGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120

Qy 121 ODVAREALGDQIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLLHCGKSFRVY 180
Db 121 ODVAREALGDQIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLLHCGKSFRVY 180

Qy 181 SNPDFIGVLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRIGALGADPATF 240
Db 181 SNPDFIGVLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRIGALGADPATF 240

Qy 241 MGMAGLDLVLTCCTENQSRNRRFGMLQGQMDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300
Db 241 MGMAGLDLVLTCCTENQSRNRRFGMLQGQMDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300

Qy 301 GVEMPTIEIYQVLYCGKNAREAAALTLGLRARKDERSH 339
Db 301 GVEMPTIEIYQVLYCGKNAREAAALTLGLRARKDERSH 339

RESULT 4
AB0975
glycerol-3-phosphate dehydrogenase (NAD) (EC 1.1.1.8) - Salmonella enterica subsp. enter
C;Species: Salmonella enterica subsp. enterica serovar Typhi
A;Note: this species has also been called Salmonella typhi
C;Date: 09-Nov-2001 #sequence_revision 09-Nov-2001 #text_change 16-Aug-2004
C;Accession: AB0975
R;Parkhill, J.; Dougan, G.; James, K.D.; Thomson, N.R.; Pickard, D.; Wain, J.; Churcher,
th, T.; Connerton, P.; Cronin, A.; Davis, P.; Davies, R.M.; Dowd, L.; White, N.; Farrar,
, S.; Moule, S.; O'Gaora, P.
Nature 413, 848-852, 2001
A;Authors: Parry, C.; Quail, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.;
A;Title: Complete genome sequence of a multiple drug resistant Salmonella enterica serova
A;Reference number: AB0502; MUID:21534947; PMID:11677608
A;Accession: AB0975
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-339 <PAR>
A;Cross-references: GB:AL513382; PIDN:CAD03294.1; PID:G16504915; GSPDB:GN00176
C;Genetics:
A;Gene: gpsA
C;Superfamily: Glycerol-3-phosphate dehydrogenase (NAD)
C;Keywords: oxidoreductase

Query Match 95.2%; Score 1640; DB 2; Length 339;
Best Local Similarity 94.7%; Pred. No. 7.3e-113;
Matches 321; Conservative 8; Mismatches 10; Indels 0; Gaps 0;

Qy 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60
Db 1 MNQRNASMTVIGAGSYGTALAITLARNGHQVVLWGHDPKHIATLEHRCNVAFPLDPVPFP 60

Qy 61 DTLHLESDLATALAASRNILVVPSPHVFGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
Db 61 DTLHLESDLATALAASRNILVVPSPHVFSDVLRQIKPLMRPDARLVWATKGLEAETGRLL 120

Qy 121 ODVAREALGDQIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLLHCGKSFRVY 180
Db 121 ODVAREALGDQIPLAVISGPTFAKELAAAGLPTAISLASTDETETFDADDLQQLLHCGKSFRVY 180

Db	68	NLAQAMEYSQDILIVVPSHAFGEILIKIKPHLKAHRLIWATKGLERNTGRLLTQTVVEEQ	127
QY	128	LGDQIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLLHCCKGSFRVYSNPDFIG	187
Db	128	LGTQYPLAVLSGPTFAKELAQGLPSAITLAANNEQFAREFQSRHCKSGFRVYINSDMTG	187
QY	188	VOLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATFMGMAGLG	247
Db	188	VOLGGAIKNVIAIGAGISDGMGFGANARTALITRGIAEITRLGISLGANTNTFMGMSGLG	247
QY	248	DLVLTCTENQSRNRRFGMMLQGQMDVQSAQEKIGQVVEGYRNTKEVRELAHRFGVEMPIT	307
Db	248	DLVLTCTDNQSRNRRFGLMLGKGLDAQWAMENIGQVVEGYNTKEAYLLAQRQGVEMPIT	307
QY	308	EEIYQVLYCGKNAREAAALTLLGRARKDE	335
Db	308	EQIYQMLFCGKSAQDVAISLLGRACKGE	335
RESULT 8			
H82637			
glycerol-3-phosphate dehydrogenase XF1802 [imported] - Xylella fastidiosa (strain 9a5c)			
C;Species: Xylella fastidiosa			
C;Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 09-Jul-2004			
C;Accession: H82637			
R;anonymous, The Xylella fastidiosa Consortium of the Organization for Nucleotide Sequen			
Nature 406, 151-157, 2000			
A;Title: The genome sequence of the plant pathogen Xylella fastidiosa.			
A;Reference number: A82515; MUID:20365717; PMID:10910347			
A;Note: for a complete list of authors see reference number A59328 below			
A;Accession: H82637			
A;Status: preliminary			
A;Molecule type: DNA			
A;Residues: 1-346 <SIM>			
A;Cross-references: UNIPROT:Q9PCH7; GB:AE004001; GB:AE003849; NID:g9106864; PIDN:AAF8461			
A;Experimental source: strain 9a5c			
R;Simpson, A.J.G.; Reinach, F.C.; Arruda, P.; Abreu, F.A.; Acencio, M.; Alvarenga, R.; A			
Briones, M.R.S.; Bueno, M.R.P.; Camargo, A.A.; Camargo, L.E.A.; Carraro, D.M.; Carrer, H			
as-Neto, E.; Docena, C.; El-Dorry, H.; Facincani, A.P.; Ferreira, A.J.S.			
Submitted to GenBank, June 2000			
A;Authors: Ferreira, V.C.A.; Ferro, J.A.; Fraga, J.S.; Franca, S.C.; Franco, M.C.; Frohm			
J.D.; Junqueira, M.L.; Kemper, E.L.; Kitajima, J.P.; Krieger, J.E.; Kuramae, E.E.; Laigh			
chado, M.A.; Madeira, A.M.B.N.; Madeira, H.M.F.; Marino, C.L.; Marques, M.V.; Martins, E			
A;Authors: Martins, E.M.F.; Matsukuma, A.Y.; Menck, C.F.M.; Miracca, E.C.; Miyaki, C.Y.;			
, F.G.; Nunes, L.R.; Oliveira, M.A.; de Oliveira, M.C.; de Oliveira, R.C.; Palmieri, D.A			
Rodrigues, V.; Rosa, A.J. de M.; de Rosa Jr., V.E.; de Sa, R.G.; Santelli, R.V.; Sawasak			
A;Authors: da Silva, A.C.R.; da Silva, F.R.; da Silva, A.M.; Silva Jr., W.A.; da Silveir			
M.; Tshako, M.H.; Vallada, H.; Van Sluys, M.A.; Verjovski-Almeida, S.; Vettore, A.L.; Z			
A;Reference number: A59328			
A;Contents: annotation			
C;Genetics:			
A;Gene: XF1802			
C;Superfamily: glycerol-3-phosphate dehydrogenase (NAD)			
Query Match 43.6%; Score 751.5; DB 2; Length 346;			
Best Local Similarity 46.7%; Pred. No. 1e-47;			
Matches 156; Conservative 54; Mismatches 111; Indels 13; Gaps 3;			
QY	8	MTVIGAGSYGTALAITLARNGHEVVLWGHDPEHIATLERDRCNAAFLPDVFPDPTLHLES	67
Db	8	I AVLGAGSWGTTAALVARHAYPTILWGRDVGIQSIDIQRNFRYLP SIMLPQT LRATT	67
QY	68	DLATALAASRNILVVVPSHVFEVLRLQIKPLMRPDARLVWATKGLEAETGRLLQDVAREA	127
Db	68	DLAAAVSGADWLVAVPSYAFTE TLRLAPLLSTGVGVAWATKGFEPGSGRFLHEVAREI	127
QY	128	LGDQIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLLHCCKGSFRVYSNPDFIG	187
Db	128	LGGDAPLAVVTGPSFAKEVTGLPTAVTVHGEYARFTQMVANMH-GPMFRAYTGN DVI G	186
QY	188	VOLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATFMGMAGLG	247

Db	187	AELGGAMKNVLAIVAIGVADGMQLGMNARAGLITRGLNEMRLS AVIGARPETLMGLAGLG	246
QY	248	DLVLTCTENQSRNRRFGMMLQGQMDVQSAQEKIGQVVEGYRNTKEVRELAHRFGVEMPIT	307
Db	247	DLVLTCTGDL SRNRLGFALGRGQSLSDAIREIGQVVESVQTSDEVMRQAEQHGVELPIS	306
QY	308	EEIYQVL-----YCGKNAREAAALTLLGRARKDE	335
Db	307	EAVRAVLREBITPYAGMKA-----LLAREQKPE	334
RESULT 9			
C97111			
glycerol 3-phosphate dehydrogenase [imported] - Clostridium acetobutylicum			
C;Species: Clostridium acetobutylicum			
C;Date: 14-Sep-2001 #sequence_revision 14-Sep-2001 #text_change 16-Aug-2004			
C;Accession: C97111			
R;Nolling, J.; Breton, G.; Omelchenko, M.V.; Markarova, K.S.; Zeng, Q.; Gibson, R.; Lee,			
.; Daly, M.J.; Bennett, G.N.; Koonin, E.V.; Smith, D.R.			
J. Bacteriol. 183, 4823-4838, 2001			
A;Title: Genome Sequence and Comparative Analysis of the Solvent-Producing Bacterium Clo			
A;Reference number: A96900; MUID:21359325; PMID:21359325			
A;Accession: C97111			
A;Status: preliminary			
A;Molecule type: DNA			
A;Residues: 1-332 <KUR>			
A;Cross-references: UNIPROT:Q97ID6; GB:AE001437; PIDN:AAK79678.1; PID:g15024677; GSPDB:G			
A;Experimental source: Clostridium acetobutylicum ATCC824			
C;Genetics:			
A;Gene: CAC1712			
C;Superfamily: Glycerol-3-phosphate dehydrogenase (NAD)			
Query Match 40.8%; Score 703; DB 2; Length 332;			
Best Local Similarity 42.1%; Pred. No. 3.5e-44;			
Matches 138; Conservative 71; Mismatches 117; Indels 2; Gaps 2;			
QY	8	MTVIGAGSYGTALAITLARNGHEVVLWGHDPEHIATLERDRCNAAFLPDVFPDPTLHLES	67
Db	4	VTFIGGSGFTALAIMLAKKHNVVWDRNKEILEDINTLRTNTRVLPNNIIPCCVKAVD	63
QY	68	DLATALAASRNILVVVPSHVFEVLRLQIKPLMRPDARLVWATKGLEAETGRLLQDVAREA	127
Db	64	DIEAKAKESKYIVLAVPSFAIREVCRKIKGFLREDQIIISIAKMBEETKKRLSEVKEE	123
QY	128	LGDQIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLLHCCKGSFRVYSNPDFIG	187
Db	124	LYKN-PVVVLSGSPSHAEVANDIPTVVVTSTDMKYAEVQDVF-MTNSFRVYTNSDIVG	181
QY	188	VOLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATFMGMAGLG	247
Db	182	VEIGGAVKNIIALASGIDGIGYGDNTKAAALTRGMSEIMRIGVKLGKPKETFFGLTGMG	241
QY	248	DLVLTCTENQSRNRRFGMMLQGQMDVQSAQEKIGQVVEGYRNTKEVRELAHRFGVEMPIT	307
Db	242	DLIVTCTSMHSRNRKAGILIGRGMSCREACDKIGMVVEGVKACHTFYELKESLGVSMPI T	301
QY	308	EEIYQVLYCGKNAREAAALTLLGRARKDE	335
Db	302	TSLYKVLFFENGDPKKEVYELMARDKONE	329
RESULT 10			
AH1316			
NAD(P)H-dependent glycerol-3-phosphate dehydrogenase homolog gpsA [imported] - Listeria			
C;Species: Listeria monocytogenes			
C;Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 16-Aug-2004			
C;Accession: AH1316			
R;Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloecker			
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi, H.			
D.; Jones, L.M.; Karst, U.			
Science 294, 849-852, 2001			
A;Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkat, G.; Madueno, E.; Maitournam, A.; Ma			
ok, C.; Schlueter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehland,			

QY	246	LGDLVLVTCTENQSRNRRFGMMLQGMDVQSAQEKIGQVVEGYRNTKEVRELAHRFGVEMP	305
Db	243	VGDLIVTCTSVHSRNRWRAGNLLGKGKLEDVLEEMGMVVEGVRTTKAAYQLSKYDVKMP	302
QY	306	ITEEIQVLYCGKNAREEAALTLLGRARKDE	335
Db	303	ITEALHQVLFGQKVETAVESLMARGKTHE	332
RESULT 13			
H83854			
NAD(P)H-dependent glycerol-3-phosphate dehydrogenase gpsA [imported] - Bacillus halodurans			
C;Species: Bacillus halodurans			
C;Date: 01-Dec-2000 #sequence_revision 01-Dec-2000 #text_change 09-Jul-2004			
C;Accession: H83854			
R;Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; Hira			
Nucleic Acids Res. 28, 4317-4331, 2000			
A;Title: Complete genome sequence of the alkaliphilic bacterium Bacillus halodurans and			
A;Reference number: A83650; MUID:20512582; PMID:11058132			
A;Accession: H83854			
A;Status: preliminary			
A;Molecule type: DNA			
A;Residues: 1-345 <STO>			
A;Cross-references: UNIPROT:Q9KCD2; GB:AP001512; GB:BA000004; NID:g10174030; PIDN:BAB053			
A;Experimental source: strain C-125			
C;Genetics:			
A;Gene: gpsA			
C;Superfamily: glycerol-3-phosphate dehydrogenase (NAD)			
Query Match 36.6%; Score 631; DB 2; Length 345;			
Best Local Similarity 38.8%; Pred. No. 6.9e-39;			
Matches 130; Conservative 70; Mismatches 123; Indels 12; Gaps 4;			
QY	7	SMTVIGAGSYGTALAITLARNGHEVVVLWGHDPHEIATLERDRCNAAFLPDVPFPD---T	62
Db	3	NVAVIGAGSWGTSLSVLADNGHHVTLVARREEIAREINERHTNETYLPSSIVAT	62
QY	63	LHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMRPDARLVWATKGLEAETGRLLQD	122
Db	63	CSMEEIIDVEI-----IVLVPTKAIRQAVRSNDVLKWPVTIVHASKGIEPGSHLRISE	117
QY	123	VAREALGDQI--PLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFVY	180
Db	118	MIEEELEPTLLKEVVVLSGPSHAEVSLRQPTVTTVSSKSLSTTKQIQDLF-MNQQFRVY	176
QY	181	SNPDFIGVOLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLAEMSRILGAALGADPATF	240
Db	177	TNEDLIGVEIGGALKNIIALACGLTNGLYGDNTKAAIMTRGLAEIGRLGVKLGASPLTF	236
QY	241	MGMAGLDLVLTCENQSRNRRFGMMLQGMDVQSAQEKIGQVVEGYRNTKEVRELAHRF	300
Db	237	AGLSGLGLDIIIVTCTSIHSRNRWRAGQMLGKGSFAEVEESMGMVVEGIRTTQAAHELAQKL	296
QY	301	GVEMPTIEEIQVLYCGKNAREEAALTLLGRARKDE	335
Db	297	QIEMPTISALYSVLFEGKKPEHAADLMGRVKKHE	331
RESULT 14			
E81953			
glycerol-3-phosphate dehydrogenase (NAD) (EC 1.1.1.8) NMA0375 [imported] - Neisseria meningitidis			
C;Species: Neisseria meningitidis			
C;Date: 05-May-2000 #sequence_revision 05-May-2000 #text_change 16-Aug-2004			
C;Accession: E81953			
R;Parkhill, J.; Achtman, M.; James, K.D.; Bentley, S.D.; Churcher, C.; Klee, S.R.; Morel			
; Holroyd, S.; Jagels, K.; Leather, S.; Moule, S.; Mungall, K.; Quail, M.A.; Rajandream,			
Nature 404, 502-506, 2000			
A;Title: Complete DNA sequence of a serogroup A strain of Neisseria meningitidis Z2491.			
A;Reference number: A81775; MUID:20222556; PMID:10761919			
A;Accession: E81953			
A;Status: preliminary			
A;Molecule type: DNA			

A;Residues: 1-329 <PAR>			
A;Cross-references: UNIPROT:Q9JWH0; GB:AL162753; GB:AL157959; NID:g7379120; PIDN:CAB8367			
A;Experimental source: serogroup A, strain Z2491			
C;Genetics:			
A;Gene: gpsA; NMA0375			
C;Superfamily: Glycerol-3-phosphate dehydrogenase (NAD)			
C;Keywords: oxidoreductase			
Query Match 36.3%; Score 624.5; DB 2; Length 329;			
Best Local Similarity 42.0%; Pred. No. 1.9e-38;			
Matches 140; Conservative 68; Mismatches 114; Indels 11; Gaps 4;			
QY	8	MTVIGAGSYGTALAITLARNGHEVVVLWGHDPHEIATLERDRCNAAFLPDVPFPDTLHLES	67
Db	3	ITVIGAGSWGTSALAHFSDQHGNRVSLWTRNADQVRMQEARENKRLPGFSFPETLEVCA	62
QY	68	DLATALAASRNILVVVPSHVFG-----EVLRQIKPLMRPDARLVWATKGLEAETGRLLQD	122
Db	63	DLADALKDSGLVLIV--TSVAGLRSSAELLKQYGAGHLP---VLAACKGFQD TGLLTFQ	117
QY	123	VAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFVYSN	182
Db	118	VLKEVLPDNKKIGVLSGPSFAQELAKQLPCAVVLASENQEWVEELVPQLNT-SVMRLYGS	176
QY	183	PDFIGVOLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLAEMSRILGAALGADPATFMG	242
Db	177	TDVIGVAVGGAVKNVMAIATGLSDGLEYNARAALVTRGLAEITRLASAMGAQPXTMVG	236
QY	243	MAGLDLVLTCENQSRNRRFGMMLQGMDVQSAQEKIGQVVEGYRNTKEVRELAHRFGV	302
Db	237	LAGIGDLIIITCTGALSRNRRVGLGLAEGKELHQVLVEIGHVSEGVSTIEEVEVENTACKYQI	296
QY	303	EMPTIEEIQVLYCGKNAREEAALTLLGRARKDE	335
Db	297	DMPITQLLQIRKEMTPQQVVERLMERSARFE	329
RESULT 15			
T35643			
glycerol-3-phosphate dehydrogenase - Streptomyces coelicolor			
C;Species: Streptomyces coelicolor			
C;Date: 05-Nov-1999 #sequence_revision 05-Nov-1999 #text_change 09-Jul-2004			
C;Accession: T35643			
R;Murphy, I.; Harris, D.; Bentley, S.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.			
submitted to the EMBL Data Library, December 1998			
A;Reference number: Z21585			
A;Accession: T35643			
A;Status: preliminary; translated from GB/EMBL/DDBJ			
A;Molecule type: DNA			
A;Residues: 1-336 <MUR>			
A;Cross-references: UNIPROT:Q9ZBS0; EMBL:AL034447; PIDN:CAA22402.1; GSPDB:GN00070; SCOEDB:			
A;Experimental source: strain A3(2)			
C;Genetics:			
A;Gene: SCOEDB:SC7Al.03			
C;Superfamily: glycerol-3-phosphate dehydrogenase (NAD)			
Query Match 36.1%; Score 621; DB 2; Length 336;			
Best Local Similarity 41.9%; Pred. No. 3.6e-38;			
Matches 139; Conservative 49; Mismatches 134; Indels 10; Gaps 3;			
QY	10	VIGAGSYGTALAITLARNGHEVVVLWGHDPHEIATLERDRCNAAFLPDVPFPDTLHLES	69
Db	9	VFGTGSWGTAFTGTVLADAGCEVTLWGRRAALADAVNSTRTNPDYLPGVLPENLRATTTA	68
QY	70	ATALAASRNILVVVPSHVFGVLRQIKPLMRPDARLVWATKGLEAETGRLLQDVAREALG	129
Db	69	AEAARDADFTVLAVPSQTLRAGLADWTPLAPGTVLVSLMKVGLGSAMRMSEV---IG	124
QY	130	DQIP-----LAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFVYSNPD	184
Db	125	DVAKVGAEERIAVVTGPNLAREIARMPAAAVVACPDETVAQRLQAACHT-PYFRPYTNTD	183
QY	185	FIGVOLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLAEMSRILGAALGADPATFMGMA	244

Db	184	VVGCELGAVKNVIGLAVGIADGMGLGDNAGSLITRGLAETTRLGVALGADPLTFSGLA	243
QY	245	GLGDLVLTCTENQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHREFGVEM	304
Db	244	GLGDLVATCSSPLSRNHTFGTNLGKGMTLEETNAVTKQTAEGVKSCESVLDLARRHGVDM	303
QY	305	PITEIYQVLYCGKNAREAAALTLLGRARKDER	336
Db	304	PITETVVAIVHEGKSPVVAVKELMSRSAPKPER	335

Search completed: April 27, 2005, 11:03:28
Job time : 41 secs

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PD 29-MAR-2001.
XX
PF 21-SEP-2000; 2000WO-CA001096.
XX
PR 22-SEP-1999; 99US-0155133P.
XX
PA (CANA) NAT RES COUNCIL CANADA.
XX
PI Zou J, Wei Y, Periappuram C, Selvaraj G, Datla R;
XX
DR WPI; 2001-257996/26.
DR P-PSDB; AAB62189.
XX
PT Manipulating glycerol-3-phosphate metabolism of plant for enhancing
PT stress tolerance, altering fatty acid content in glycerolipids, by
PT expressing in plant feedback defective glycerol-3-phosphate dehydrogenase
PT gene.
XX
PS Claim 5; Fig 1; 39pp; English.
XX
CC The invention provides a method for genetically transforming a plant so
CC that it expresses a heterologous glycerol-3-phosphate dehydrogenase
CC (G3PD) that is less sensitive to feedback inhibition than wild-type G3PD.
CC The method involves providing a vector comprising a DNA sequence encoding
CC G3PD that is less sensitive to feedback inhibition than wild-type G3PD
CC and transforming the plant with the vector. The method is useful for
CC expressing a heterologous G3PD less sensitive to feedback inhibition than
CC wild-type G3PD in an oil seed bearing plant, such as Arabidopsis thaliana
CC or Brassica. The vectors are useful for producing a genetically altered
CC plant having altered fatty acid content in its glycerolipids, especially
CC elevated levels of C16 fatty acids and increased osmotic stress tolerance
CC relative to the wild type. The present sequence represents the DNA
CC encoding the E. coli gpsA2FR protein. The gene gpsA2FR is an allele of
CC the E. coli gpsA gene, and encodes an altered version of the GPDH protein
CC defective in feedback inhibition. This gpsA2FR gene can be used in the
CC vectors and method of the invention
XX
SQ Sequence 1020 BP; 214 A; 274 C; 304 G; 228 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 3.46e-158 Length: 1020
Score: 1722.00 Matches: 339
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 5 Gaps: 0

US-10-088-079-2 (1-339) x AAF57428 (1-1020)

QY 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20
Db 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCCGGCTCGTACGGCACCGCTCTT 60

QY 21 AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis 40
Db 61 GCCATCACCCCTGGCAAGAAATGGCCACGAGGTTCTCTCTGGGGCCATGACCTGAACAT 120

QY 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProphePro 60
Db 121 ATCGCAACGCTTGAACGGACCGCTGTAAACGCCGGTTTCTCCCCGATGTGCCTTTTCCC 180

QY 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80
Db 181 GATACGCTCCATCTTGAAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAATATCTC 240

QY 81 ValValValProSerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArg 100
Db 241 GTCGTCGTACCCAGCCATGTCTTTGGTGAAGTGTGCGCCAGATTAAACCACTGATGCGT 300

QY 101 ProAspAlaArgLeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120
Db 301 CCTGATCGCGCTCTGGTGTGGGCGGACCAAGGGCTGGAAGCGGAACCGGACGCTCTGTTA 360

QY 121 GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140
Db 361 CAGGACGTGGCCGTGAGGCCCTTAGCGATCAAAATTCCGCTGGCGGTATCTCTGGCCCA 420

QY 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160
Db 421 ACGTTTGCAGAAAGAACTGGCGGCAGGTTTACCGACAGCTATTTCGCTGGCCTCGACCGAT 480

QY 161 GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180
Db 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGTGCACTGCGGCAAAAGTTTCCGCGTTTAC 540

QY 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle 200
Db 541 AGCAATCCGGATTTCATTGGCGTGCAGCTTGGCGGCGGTGAAACGTTATTGCCATT 600

QY 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220
Db 601 GGTGGGGGATGTCCGACGGTATCGGTTTGGTGCGAATGCGCGTACGGCGCTGATCACC 660

QY 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240
Db 661 CGTGGGCTGGCTGAAATGTCCGCTTGGTGCGGCGCTGGGTGCCGACCTGCCACCTTT 720

QY 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260
Db 721 ATGGGCATGGCGGGCTTGGCGATCTGGTCTTACCTGTACCGAAAACAGTCCGCGTAAC 780

QY 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280
Db 781 CGCCGTTTGGCATGATGTCGGTCAGGCGATGGATGTACAAAGCGCGCAGGAGAAGATT 840

QY 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300
Db 841 GGTGAGTGTGGAAGGCTACCGCAATACGAAAGAGTCCCGCAACTGGCGCATCGCTTC 900

QY 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320
Db 901 GGCCTTGAATGCCAATAACCGAGGAATTATCAAGTATTATTTGCGGAAAAACGCG 960

QY 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSerHis 339
Db 961 CGCAGGCAGCATTGACTTTACTAGGTCGTGCACGCAAGGACGAGCGCAGCAGCCAC 1017

RESULT 2
AAS52655
ID AAS52655 standard; DNA; 1020 BP.
XX
AC AAS52655;
XX
DT 13-FEB-2002 (first entry)
XX
DE E. coli DNA for cellular proliferation protein #377.
XX
KW Antisense; ds; prokaryotic cellular proliferation gene; antibiotic;
KW antibacterial; drug design.
XX
OS Escherichia coli.
XX
PN WO200170955-A2.
XX
PD 27-SEP-2001.
XX
PF 21-MAR-2001; 2001WO-US0009180.
XX
PR 21-MAR-2000; 2000US-0191078P.
PR 23-MAY-2000; 2000US-0206848P.
PR 26-MAY-2000; 2000US-0207727P.
PR 23-OCT-2000; 2000US-0242578P.
PR 27-NOV-2000; 2000US-0253625P.
PR 22-DEC-2000; 2000US-0257931P.
PR 16-FEB-2001; 2001US-0269308P.
XX

PA (ELIT-) ELITRA PHARM INC.
XX
PI Haselbeck R, Ohlsen KL, Zyskind JW, Wall D, Trawick JD, Carr GJ;
PI Yamamoto RT, Xu HH;
XX
DR WPI; 2001-611495/70.
DR P-PSDB; AAU34796.
XX
XX New polynucleotides for the identification and development of
PT antibiotics, comprise sequences of antisense nucleic acids.
PT
XX
PS Claim 27; SEQ ID NO 6292; 511pp; English.
XX
CC The invention relates to antisense inhibitors of genes essential to
CC prokaryotic cellular proliferation, their use in identifying the genes,
CC their use in the discovery of novel antibiotics, the essential genes
CC themselves and the encoded proteins. The prokaryotes used are Escherichia
CC coli, Staphylococcus aureus, Salmonella typhi, Klebsiella pneumoniae,
CC Pseudomonas aeruginosa and Enterococcus faecalis. The invention is also
CC useful for the identification of potential new targets for antibiotic
CC development. The antisense nucleic acids can also be used to identify
CC proteins used in proliferation, to express these proteins, and to obtain
CC antibodies capable of binding to the expressed proteins. The proteins can
CC be used to screen compounds in rational drug discovery programmes. The
CC antisense nucleic acid sequence is also useful to screen for homologous
CC nucleic acids which are required for cell proliferation in a wide variety
CC of organisms. The present sequence encodes an essential prokaryotic
CC cellular proliferation protein. Note: The sequence data for this patent
CC did not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 1020 BP; 213 A; 275 C; 304 G; 228 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 6.78e-158 Length: 1020
Score: 1719.00 Matches: 338
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 99.71% Mismatches: 0
Query Match: 99.83% Indels: 0
DB: 4 Gaps: 0

US-10-088-079-2 (1-339) x AAS52655 (1-1020)

Qy 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20
Db 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCGCGCTCGTACGGCACCGCTCTT 60
Qy 21 AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis 40
Db 61 GCCATCACCTGGCAAGAAATGSCCAGAGGTTGTCTCTGGGCCCATGACCCCTGAACAT 120
Qy 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProPhePro 60
Db 121 ATCGCAACGCTTGAACGCGACCCGCTGTAAACGCCCGCTTCTCCCCGATGTGCCTTTCCC 180
Qy 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80
Db 181 GATACGCTCCATCTTGAAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAATAFTCTC 240
Qy 81 ValValValProSerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArg 100
Db 241 GTCGTCGTACCCAGCCATGCTTTGGTGAAGTGTCTGCGCCAGATTAAACCACTGATCGGT 300
Qy 101 ProAspAlaArgLeuValTipAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120
Db 301 CCTGATGCGGCTCTGGTGTGGGCGACCAAGGGCTGGAAGCGGAACCGGACGTCGTGTTA 360
Qy 121 GlnAspValAlaAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140
Db 361 CAGGACGTGGCGGTGAGGCCCTTAGCGGATCAAATTCGCTGGCGGTTATCTCTGGCCCA 420
Qy 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160

Db 421 ACGTTTGCGAAGAAGAACTGGCGGCAGGTTTACCGACAGCTATTTCGCTGGCTCGACCGAT 480
Qy 161 GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180
Db 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACACTGGGGCAAAAGTTTCCGCGTTAC 540
Qy 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle 200
Db 541 AGCAATCCGGATTTCAATTGGCGTGACGCTTGGCGGCGCGGTGAAAAACGTTATTGCCATT 600
Qy 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220
Db 601 GGTGCGGGGATGTCGACGGTATCGGTTTGGTGCGAATGCGCGTACGGCGTGATCACC 660
Qy 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240
Db 661 CGTGGGCTGGCTGAAATGTGCGGTCTTTGGTGCGGCGCTGGGTGCCGACCTGCCACCTTT 720
Qy 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260
Db 721 ATGGCATGGCGGGCTTGGCGATCTGGTGCTTACCTGTACCGACAACCACTCGCGTAAC 780
Qy 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280
Db 781 CGCGGTTTTGGCATGATGCTCGGTGAGGGCATGGATGTACAAAGCGCGCAGGAGAGATT 840
Qy 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300
Db 841 GGTGAGGTGGTGAAGGCTACCGCAATACGAAAGAAAGTCCCGGAACCTGGCGCATCGCTTC 900
Qy 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320
Db 901 GCGGTTGAAATGCCAATAACCGAGGAAATTTATCAAGTATTATATTCGGAAAAAACGCG 960
Qy 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSerHis 339
Db 961 CGCGAGGCAGCATTTGACTTTACTAGGTGTCGACCGCAAGGACGAGCGCAGCAGCCAC 1017
RESULT 3
ACA32689
ID ACA32689 standard; DNA; 1020 BP.
XX
AC ACA32689;
XX
DT 19-JUN-2003 (first entry)
XX
DE Prokaryotic essential gene #14346.
XX
KW Antisense; ds; prokaryotic essential gene; cell proliferation;
KW drug design; gene.
XX
OS Escherichia coli.
XX
PN WO200277183-A2.
XX
PD 03-OCT-2002.
XX
PF 21-MAR-2002; 2002WO-US0009107.
XX
PR 21-MAR-2001; 2001US-00815242.
PR 06-SEP-2001; 2001US-00948993.
PR 25-OCT-2001; 2001US-0342923P.
PR 08-FEB-2002; 2002US-00072851.
PR 06-MAR-2002; 2002US-0362699P.
XX
PA (ELIT-) ELITRA PHARM INC.
XX
PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
XX
DR WPI; 2003-029926/02.
DR P-PSDB; ABU28819.

XX New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.
XX
PS Claim 14; SEQ ID NO 20559; 1766pp; English.
XX
CC The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
CC prokaryotic essential genes. Note: The sequence data for this patent did
CC not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 1020 BP; 213 A; 275 C; 304 G; 228 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 6.78e-158 Length: 1020
Score: 1719.00 Matches: 338
Percent Similarity: 100.00% Conservatives: 1
Best Local Similarity: 99.71% Mismatches: 0
Query Match: 99.83% Indels: 0
DB: 8 Gaps: 0

US-10-088-079-2 (1-339) x ACA332689 (1-1020)

Qy 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20
Db 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCGGCTCGTACGGCACCGCTCTT 60
Qy 21 AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis 40
Db 61 GCCATCACCCCTGGCAAGAAATGGCCACGAGGTTGTCTCTGGGGCCATGACCCCTGAACAT 120
Qy 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProphePro 60
Db 121 ATCGCAACGCTTGAACGCGACCGCTGTAAACGCGGTTTCTCCCCGATGTGCTTTTCCC 180
Qy 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80
Db 181 GATACGCTCCATCTTGAAGCGATCTCGCCACTCGCTGGCAGCCAGCCGTAATATTCTC 240
Qy 81 ValValValProSerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArg 100
Db 241 GTCGTGTCACCCAGCCATGCTTTTGGTGAAGTGTCTGGCCAGATTAAACCACTGATCGGT 300
Qy 101 ProAspAlaArgLeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120

Db 301 CCTGATGCGCGTCTGGTGTGGGCGACCAAGGGCTGGAAGCGGAAACCGGACGTCTGTTA 360
Qy 121 GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140
Db 361 CAGGACGTGGCGGTGAGGCCTTAGGCGATCAAAATTCCGCTGGCGGTTATCTCTGGCCCCA 420
Qy 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160
Db 421 ACGTTTGCAGAAAGAACTGGCGGAGGTTTACCGACAGCTATTTCGTGGCTCGACCCGAT 480
Qy 161 GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180
Db 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGCGGCAAAAGTTTCCGCGTTTAC 540
Qy 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle 200
Db 541 AGCAATCCGGATTTCATTGGCGTGCAGCTTGGCGGCGGTTGAAACGTTATTGCCATT 600
Qy 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220
Db 601 GGTGCGGGGATGTCGACGGTATCGGTTTGGTGCGAATCGCGTACGGCGCTGATCACC 660
Qy 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240
Db 661 CGTGGGCTGGCTGAAATGTCCGCTCTTGGTGGCGGCTGGGTGCCGACCCCTGCCACCTTT 720
Qy 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260
Db 721 ATGGGCATGGCGGGCTTGGCGATCTGGCTTACCTGTACCGACACACAGTCGCGTAAC 780
Qy 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280
Db 781 CGCCGTTTTGGCATGATGTCGGTCAAGGCGATGGATGTACAAAGCGCGCAGGAGAAGATT 840
Qy 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300
Db 841 GGTCAAGTGGTGAAGGCTACCGCAATACGAAAGAGTCCCGCAACTGGCGCATCGCTTC 900
Qy 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320
Db 901 GGCCTTGAATGCCAATAACCGAGGAATTTATCAAGTATTATTTGCGGAAAAAACGCG 960
Qy 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSerHis 339
Db 961 CGCGAGGCAGCATTGACTTTACTAGGTCGTGCACGCAAGGACGAGCGCAGCAGCCAC 1017
RESULT 4
ADT48853
ID ADT48853 standard; cDNA; 1020 BP.
XX
AC ADT48853;
XX
DT 02-DEC-2004 (first entry)
XX
DE Bacterial polynucleotide #23604.
XX
KW Recombinant DNA construct; transformed plant; improved plant property;
KW cold tolerance; heat tolerance; drought tolerance; herbicide; osmosis;
KW pathogen tolerance; pest tolerance; plant disease resistance;
KW cell cycle pathway modification; plant growth regulator;
KW homologous recombination; seed oil yield; protein yield; carbohydrate;
KW nitrogen; phosphorus; photosynthesis; lignin; galactomannan;
KW bacterial polynucleotide; gene; ss.
OS Bacteria.
XX
XX US2003233675-A1.
PN
XX
XX 18-DEC-2003.
XX
PF 20-FEB-2003; 2003US-00369493.
XX
PR 21-FEB-2002; 2002US-0360039P.

XX (CAOY/) CAO Y.
PA (HINK/) HINKLE G J.
PA (SLAT/) SLATER S C.
PA (CHEN/) CHEN X.
PA (GOLD/) GOLDMAN B S.
XX
PI Cao Y, Hinkle GJ, Slater SC, Chen X, Goldman BS;
XX WPI; 2004-061375/06.
DR
XX
XX
PT New recombinant DNA construct comprising a promoter positioned to provide
PT for expression of a polynucleotide encoding a polypeptide from a
PT microbial source, useful for producing plants with improved properties.
XX
PS Claim 1; SEQ ID NO 47291; 122pp; English.
XX
CC The invention relates to a recombinant DNA construct comprising a
CC promoter functional in a plant cell, where the promoter is positioned to
CC provide for expression of a polynucleotide encoding a polypeptide from a
CC microbial source. The invention also relates to a transformed plant
CC comprising the recombinant DNA construct and a method of producing a
CC transformed plant having an improved property. The plant is a crop plant
CC such as maize or soybean. The method of producing a transformed plant
CC having an improved property comprises transforming a plant with the
CC recombinant DNA construct and growing the transformed plant, where the
CC polynucleotide or polypeptide is useful for improving plant properties.
CC The recombinant DNA construct is useful for producing plants with
CC improved plant properties, e.g. improved cold, heat or drought tolerance,
CC tolerance to herbicides, extreme osmotic conditions, pathogens or pests,
CC increased resistance to plant disease, better growth rate by modification
CC of the cell cycle pathway with plant growth regulators, increased rate of
CC homologous recombination, modified seed oil or protein yield and/or
CC content, improved yield by modification of carbohydrate, nitrogen or
CC phosphorus use and/or uptake, by modification of photosynthesis or by
CC providing improved plant growth and development under at least one stress
CC condition, improved lignin production or improved galactomannan
CC production. This sequence represents a bacterial polynucleotide used in
CC the scope of the invention. Note: The sequence data for this patent did
CC not form part of the printed specification but was obtained in electronic
CC format from USPTO at seqdata.uspto.gov/sequence.html.

XX
SQ Sequence 1020 BP; 213 A; 275 C; 304 G; 228 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 6.78e-158 Length: 1020
Score: 1719.00 Matches: 338
Percent Similarity: 100.00% Conservative: 1
Best Local Similarity: 99.71% Mismatches: 0
Query Match: 99.83% Indels: 0
DB: 13 Gaps: 0

US-10-088-079-2 (1-339) x ADT48853 (1-1020)

Qy 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20
Db 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCCGGCTCGTACGGCACCCGCTCTT 60
Qy 21 AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTyrGlyHisAspProGluHis 40
Db 61 GCCATCACCTGGCAAGAAATGGCCACGAGGTTGCTCTCGGGCCATGACCCCTGAACAT 120
Qy 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProPhePro 60
Db 121 ATCGCAACGCTTGAAACGCGACCGCTGTAAACGCCGCGTTCTCTCCCGATGTGCTTTTCCC 180
Qy 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80
Db 181 GATACGCTCCATCTTGAAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAATATCTC 240
Qy 81 ValValValProSerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArg 100
Db 241 GTCGTCGTACCCAGCCATGCTCTTTGGTGAAGTGTCTGCCAGTGTCTGCCAGATTAACCACTGATGCGT 300

Qy 101 ProAspAlaArgLeuValTrrAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120
Db 301 CCTGATGCGCGTCTGGTGTGGCGACCAAGGGCTGGAAGCGGAACCGGACGTCGTGTTA 360
Qy 121 GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140
Db 361 CAGGACGTGGCGGTGAGGCTTAGGCGATCAAATTCGCTGGCGTTATCTCTGGCCCA 420
Qy 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160
Db 421 ACGTTTGCAGAAAGAACTGGCGGCAGGTTTACCGACAGCTATTTCGCTGGCTCGACCGAT 480
Qy 161 GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180
Db 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGCGGCAAAAGTTTCCGCGTTTAC 540
Qy 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle 200
Db 541 AGCAATCCGGATTTTCATTGGCGTGACGCTTGGCGGCGCGGTGAAAAACGTTATTGCCATT 600
Qy 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220
Db 601 GGTGCGGGGATGTCGACGGTATCGGTTTTTGGTGCNAATGCGCGTACGGCGCTGATCAC 660
Qy 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240
Db 661 CGTGGGCTGGCTGAAATGTCCGCTCTTGGTGCGGCGCTGGGTGCCGACCCCTGCCACTTT 720
Qy 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260
Db 721 ATGGCATGGCGGGGCTTGGCGATCTGGTGTCTTACCTGTACCGACAACCAACGATCGCGTAA 780
Qy 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280
Db 781 CGCCGTTTTGGCATGATGCTCGGTGAGGCGATGGATGTACAAAGCGCGCAGGAGAGATT 840
Qy 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300
Db 841 GGTGAGTGGTGAAGGCTACCGCAATACGAAAGAGTCCGCGAATCGGCGCATCGCTTC 900
Qy 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320
Db 901 GCGGTTGAAATGCCAATAACCGAGGAAATTTATCAAGTATTATATTGCGGAAAAAACGCG 960
Qy 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSerHis 339
Db 961 CGCGAGGCAGCATTTGACTTTACTAGGTGCTGTCACGCAAGGACGAGCGCAGAGCCAC 1017
RESULT 5
ACA51335
ID ACA51335 standard; DNA; 1020 BP.
XX
AC ACA51335;
XX
DT 19-JUN-2003 (first entry)
XX
DE Prokaryotic essential gene #32992.
XX
KW Antisense; ds; prokaryotic essential gene; cell proliferation;
KW drug design; gene.
XX
OS Salmomella typhi.
XX
PN WO200277183-A2.
XX
PD 03-OCT-2002.
XX
PF 21-MAR-2002; 2002WO-US0009107.
XX
PR 21-MAR-2001; 2001US-00815242.
PR 06-SEP-2001; 2001US-00948993.
PR 25-OCT-2001; 2001US-0342923P.

PR 08-FEB-2002; 2002US-00072851.
PR 06-MAR-2002; 2002US-0362699P.
XX
PA (ELIT-) ELITRA PHARM INC.
XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
XX
DR WPI; 2003-0299926/02.
DR P-PSDB; ABU47465.
XX
PT New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.
XX
PS Claim 14; SEQ ID NO 39205; 1766pp; English.
XX
CC The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
CC prokaryotic essential genes. Note: The sequence data for this patent did
CC not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 1020 BP; 217 A; 271 C; 316 G; 216 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 3.41e-150 Length: 1020
Score: 1640.00 Matches: 321
Percent Similarity: 97.05% Conservative: 8
Best Local Similarity: 94.69% Mismatches: 10
Query Match: 95.24% Indels: 0
DB: 8 Gaps: 0

US-10-088-079-2 (1-339) x ACA51335 (1-1020)

QY 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20
Db 1 ATGAACCAAGTAATGCGTCAATGACAGTCATCGGTGCGGCTCGTACGGCACCGCTCTC 60
QY 21 AlaIleThrLeuAlaArghenGlyHisGluValValLeuTrpGlyHisAspProGluHis 40
Db 61 GCCATCACTCTGGCGAGAAACGGCCACCAGGTTGCTCTGTGGGGCCACGACCAACAT 120
QY 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProPhePro 60
Db 121 ATCGGACCCCTGGAGCACGATCGCTGCAACGTCGCGTTCCTTCCCGATGTGCCTTTCCC 180

QY 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80
Db 181 GATACGTTACACCTGGAAGCGACTTAGCAACCGCGCTGGCGGCCAGTCGTAAACATTCTG 240
QY 81 ValValValProSerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArg 100
Db 241 GTGGTGGTGCCAAAGCCATGTTTTCAGCGACGTGCTCGGCAGATTAAACCGCTGATCGCT 300
QY 101 ProAspAlaArgLeuValTirAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120
Db 301 CCGGATGCGCGTCTGGTATGGGCGACCAAGGCTGGAAGCGGAAACGGGGCGCCTGTTG 360
QY 121 GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140
Db 361 CAGGATGTCGCTCGGAGGCGTTAGGCGATCAAATCCCGCTGGCGGTGATTCCGGTCCG 420
QY 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160
Db 421 ACGTTCGCTAAAGAGCTGGCGGGGGTTTGCCGACGGCAATCTCGCTAGCCTCAACCGAT 480
QY 161 GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180
Db 481 GAGACCTTTGCCGACGATCTCCAGCAACTGTTGCACCTGCGGAAAAAGTTTTCGCGTCTAT 540
QY 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle 200
Db 541 ATCAATGCGGATTTATCGGCGTGCAGCTTGGCGGCGGTGAAAAACGTGATTCGATT 600
QY 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220
Db 601 GGCGGGGGATGCTGACGGCATCGGCTTCGGCGCGAACCCTCCGCGCTAATCACG 660
QY 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240
Db 661 CGTGGACTGACCGAAATGTGCGGGCTTGGCGCAGCGCTTGGTGCAGTCCCGCACCTTT 720
QY 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260
Db 721 ATGGGGATGGCGGGTTTAGCGGATCTGGTCTGACCTGTACCGACAACCATGCGCGCAAC 780
QY 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280
Db 781 CGTCGTTTTGGCATGATGCTTGGCCAGGGCATGGACGTTAAAGGCGCGCAGGATAAGATT 840
QY 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300
Db 841 GGCCAGGTGTCGAAAGGCTATCGCAATACGAAAGAAAGTTTCGTGAATTGGCGCACCGTTT 900
QY 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320
Db 901 GGTGTTGAATGCCAATAACCGAGGAAATTTATCAAGTATTGTATTGCGGAAAAAACGCG 960
QY 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSerHis 339
Db 961 CGCGAGGCAGCATTAACGTTATTAGTTCGCGCCCGCAAGGAAGAGCTGAGTCGCCAC 1017
RESULT 6
ACA31966
ID ACA31966 standard; DNA; 1017 BP.
XX
AC ACA31966;
XX
DT 19-JUN-2003 (first entry)
XX
DE Prokaryotic essential gene #13623.
XX
KW Antisense; ds; prokaryotic essential gene; cell proliferation;
KW drug design; gene.
XX
OS Enterobacter cloacae.
XX
PN WO200277183-A2.
XX

KW Antisense; ds; prokaryotic essential gene; cell proliferation;
KW drug design; gene.
XX Klebsiella pneumoniae.
OS
XX WO200277183-A2.
PN
XX
PD 03-OCT-2002.
XX
PF 21-MAR-2002; 2002WO-US009107.
XX
PR 21-MAR-2001; 2001US-00815242.
PR 06-SEP-2001; 2001US-00948993.
PR 25-OCT-2001; 2001US-0342923P.
PR 08-FEB-2002; 2002US-00072851.
PR 06-MAR-2002; 2002US-0362699P.
XX
PA (ELIT-) ELITRA PHARM INC.
PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
XX
DR WPI; 2003-029926/02.
DR P-PSDB; ABU31968.
XX
PT New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.
XX
PS Claim 14; SEQ ID NO 23708; 1766pp; English.
XX
CC The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
CC prokaryotic essential genes. Note: The sequence data for this patent did
CC not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 1017 BP; 194 A; 306 C; 319 G; 198 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 4.98e-145 Length: 1017
Score: 1587.00 Matches: 313
Percent Similarity: 95.58% Conservative: 11
Best Local Similarity: 92.33% Mismatches: 15
Query Match: 92.16% Indels: 0
DB: 8 Gaps: 0

US-10-088-079-2 (1-339) x ACA35838 (1-1017)
QY 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20
Db 1 ATGAACGCACCTTAATGCTGCAATGACTGTGATCGGTGCCGGCTCTTACGGCACCGCTCTT 60
QY 21 AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis 40
Db 61 GCCATCACCTTGGCAAGAAATGGCCACCACGTTGTCTGTGGGGCCATGACCCGAAACAT 120
QY 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProphePro 60
Db 121 ATCGCGACGCTGCAACACGATCGCTGCAACGCCGCGTTCCTTCCCGATGTGCCTTTCCCG 180
QY 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80
Db 181 GATACGCTGCATCTTGAGAGCGACCTGGCCACCGCGCTGGCCGCCAGCCGACATCCTT 240
QY 81 ValValValProSerHisValPheGlyGluValValLeuArgGlnIleIleValProLeuMetArg 100
Db 241 GTCGTGGTGGCGAGCCATGTATTTCGGTCAGGTGTTACGCCAGATTAAACCGCTGATCGGT 300
QY 101 ProAspAlaArgLeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120
Db 301 TCCGACGCGCGCTGGTGTGGGCCACCAAGGCCCTTGAGGCCGAAACCGCGCGTCTGCTG 360
QY 121 GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140
Db 361 CAGGACGTGGCGCTGAAGCGCTGGCGCATGATATTCCGCTGGCGGTGATCTCGGGGCCA 420
QY 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160
Db 421 ACCTTCGCCAAAGAGCTGGCCGCGCTGCCGACGGCGATTTCGTGGCGGCCACGGAT 480
QY 161 GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180
Db 481 CCGCAGTTTGGCGAGGACCTTCAGCGCTTACTGCACGTGGGCAAAAGCTTCGCGTCTAC 540
QY 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle 200
Db 541 ATCAACCCGGACTTTATCGGCGTGCAGCTCGGCGCGCGGTGAAACCGTCATTGCCATC 600
QY 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220
Db 601 GGGGCAGGTATGTCGGACGGCATCGGCTTCGGCGCCCAATCGCGGTACGCGCTGATTACC 660
QY 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240
Db 661 CGTGGCTGTGTGAAATGTCCCGCTCGCGCGCGCGCTGGCGCGCGATCCGGAAACCTTT 720
QY 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260
Db 721 ATGGGCATGGCGGCGCTCGGTGACCTGGTGCTCACCTGCACCGACACCGAGTCCCGTAAC 780
QY 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280
Db 781 CGTCGCTTCGCGCATGATGCTCGGCCAGGGTATGGACGTGCAGAGCGCCAGGACAAGATT 840
QY 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300
Db 841 GGCCAGGTGGTTGAAGGCTACCGCAATACCAAGGAAGTTTCGCGCTTCGGCACAGCGTTTA 900
QY 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320
Db 901 GGTGTCGAAATGCCAATAACCGAGGAAATTATCAGGTATTGTATTGCGGAAAAAATTGCG 960
QY 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSerHis 339
Db 961 CGCGAGGCAGCATTGACCTTATTGGGTGCGGCCCGCAAGGACGAGCGCAGCAAT 1017
RESULT 8
ACH97774
ID ACH97774 standard; DNA; 1038 BP.

XX ACH97774;
AC 29-JUL-2004 (first entry)
XX Klebsiella pneumoniae polynucleotide seqid 3569.
DE
XX
XX Recombinant expression vector; transcription regulatory element;
KW Klebsiella pneumoniae protein; antibacterial; vaccine; gene; ds.
XX
XX Klebsiella pneumoniae.
OS
XX
XX US6610836-B1.
PN
XX
XX 26-AUG-2003.
PD
XX
XX 27-JAN-2000; 2000US-00489039.
PF
XX
XX 29-JAN-1999; 99US-0117747P.
PR
XX
XX (GENO-) GENOME THERAPEUTICS CORP.
PA
XX
XX Breton GL, Osborne M;
PI
XX
XX WPI; 2003-895346/82.
DR
XX P-PSDB; ABO64223.
XX
PT New nucleic acid encoding a Klebsiella pneumoniae polypeptide, useful for
PT preparing a vaccine composition against Klebsiella pneumoniae.
XX
PS Disclosure; SEQ ID NO 3569; 932pp; English.
XX
CC The invention describes a new isolated nucleic acid encoding a Klebsiella
CC pneumoniae polypeptide. Also described are: a recombinant expression
CC vector comprising the nucleic acid, operably linked to a transcription
CC regulatory element; and a cell comprising the recombinant expression
CC vector. The nucleic acid is useful for preparing a vaccine composition
CC against Klebsiella pneumoniae. This sequence encodes a Klebsiella
CC pneumoniae polypeptide of the invention
XX
SQ Sequence 1038 BP; 201 A; 310 C; 324 G; 203 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 5.12e-145 Length: 1038
Score: 1587.00 Matches: 313
Percent Similarity: 95.58% Conservative: 11
Best Local Similarity: 92.33% Mismatches: 15
Query Match: 92.16% Indels: 0
DB: 11 Gaps: 0

US-10-088-079-2 (1-339) x ACH97774 (1-1038)

QY 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20
Db 19 ATGAACGCACTTAATGCTGCAATGACTGTGATCGGTGCCGCTCTTACGGCACCGCTCTT 78

QY 21 AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis 40
Db 79 GCATCACCTTGGCAAGAAATGGCCACCACCGTTGTGCTGTGGGGCCATGACCCGAAACAT 138

QY 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProPhePro 60
Db 139 ATCGCGACGCTGCAACACGATCGCTGCAACGCCGCTTCTTCCGATGTGCCCTTCCCG 198

QY 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80
Db 199 GATACGCTGCATCTTGAGAGCGACCTGGCCACCGCGTGGCCGCCAGCCCGCATCCTT 258

QY 81 ValValValProSerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArg 100
Db 259 GTCGTGGTGCCGAGCCATGTATTCGGTCAGGTGTTACGCCAGATTAAACCGCTGATCGGT 318

QY 101 ProAspAlaArgLeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120

Db 319 TCCGACGCGCGCTGGTGTGGGCCACCAAGGCCCTTGAGGCCGAACCGCGCTGTGCTG 378
QY 121 GlnAspValAlaAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140
Db 379 CAGGACGTGGCGCGTGAAGCGCTGGGCGATGATATTCGCTGGCGGTGATCTCGGGGCCA 438
QY 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160
Db 439 ACCTTCGCCAAAGAGCTGGCGCGCGCTGCGGACGGCGATTTCGTGGCGGCCACGGAT 498
QY 161 GlnThrPheAlaAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180
Db 499 CCGCAGTTTTCGGGAGGACCTTCAGCGCCTACTGCACTGCGGCAAAAGCTTCCGCGCTAC 558
QY 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyValValLysAsnValIleAlaIle 200
Db 559 ATCAACCCCGACTTTATCGGCGTGCAGCTCGGCGCGCGGTGAAAAACGTCATTGCCATC 618
QY 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220
Db 619 GGGGCAGGTATGTCGGATGGCATCGGCTTCGGCGGCAATGCGCGTACGGCGCTGATTACC 678
QY 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240
Db 679 CGTGGGCTGGTGGAAATGTCCCGCTCGGCGCGCGCTGGGCGCGCGATCCGGAAACCTTT 738
QY 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260
Db 739 ATGGGCATGGCCGCGCTCGGTGACCTGGTGCTCACCTGCACCGACCAACGATCCCGTAAC 798
QY 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280
Db 799 CGTCGCTTCGGCATGATGCTCGGCCAGGGTATGGACGTGCAGAGCGCCAGGACAAGATT 858
QY 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300
Db 859 GGCCAGGTGTTGAAGCTACCGCAATACCAAGGAAGTTCGCGTTCCTGGCACAGCGTTTA 918
QY 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320
Db 919 GGTGTCGAAATGCCAATAACCGAGGAAATTTATCAGGTATTGTATTGGGAAAAAATTGCG 978
QY 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSerHis 339
Db 979 CGCGAGGCAGCATTGACCTTATTGGGTTCGCGCCCGCAAGGACGAGCGCAGCAAT 1035
RESULT 9
ACA49224
ID ACA49224 standard; DNA; 1023 BP.
XX
AC ACA49224;
XX
DT 19-JUN-2003 (first entry)
XX
DE Prokaryotic essential gene #30881.
XX
KW Antisense; ds; prokaryotic essential gene; cell proliferation;
KW drug design; gene.
XX
OS Salmonella paratyphi.
XX
PN WO200277183-A2.
XX
PD 03-OCT-2002.
XX
PF 21-MAR-2002; 2002WO-US009107.
XX
PR 21-MAR-2001; 2001US-00815242.
PR 06-SEP-2001; 2001US-00948993.
PR 25-OCT-2001; 2001US-0342923P.
PR 08-FEB-2002; 2002US-00072851.
PR 06-MAR-2002; 2002US-0362699P.

XX PA (ELIT-) ELITRA PHARM INC.

XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;

PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;

XX WPI; 2003-029926/02.

DR P-PSDB; ABU45354.

XX

PT New antisense nucleic acids, useful for identifying proteins or screening

PT for homologous nucleic acids required for cellular proliferation to

PT isolate candidate molecules for rational drug discovery programs.

XX

PS Claim 14; SEQ ID NO 37094; 1766pp; English.

XX

CC The invention relates to an isolated nucleic acid comprising any one of

CC the 6213 antisense sequences given in the specification where expression

CC of the nucleic acid inhibits proliferation of a cell. Also included are:

CC (1) a vector comprising a promoter operably linked to the nucleic acid

CC encoding a polypeptide whose expression is inhibited by the antisense

CC nucleic acid; (2) a host cell containing the vector; (3) an isolated

CC polypeptide or its fragment whose expression is inhibited by the

CC antisense nucleic acid; (4) an antibody capable of specifically binding

CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular

CC proliferation or the activity of a gene in an operon required for

CC proliferation; (7) identifying a compound that influences the activity of

CC the gene product or that has an activity against a biological pathway

CC required for proliferation, or that inhibits cellular proliferation; (8)

CC identifying a gene required for cellular proliferation or the biological

CC pathway in which a proliferation-required gene or its gene product lies

CC or a gene on which the test compound that inhibits proliferation of an

CC organism acts; (9) manufacturing an antibiotic; (10) profiling a

CC compound's activity; (11) a culture comprising strains in which the gene

CC product is overexpressed or underexpressed; (12) determining the extent

CC to which each of the strains is present in a culture or collection of

CC strains; or (13) identifying the target of a compound that inhibits the

CC proliferation of an organism. The antisense nucleic acids are useful for

CC identifying proteins or screening for homologous nucleic acids required

CC for cellular proliferation to isolate candidate molecules for rational

CC drug discovery programs, or for screening homologous nucleic acids

CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,

CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target

CC prokaryotic essential genes. Note: The sequence data for this patent did

CC not form part of the printed specification, but was obtained in

CC electronic format directly from WIPO at

CC ftp.wipo.int/pub/published_pct_sequences

XX

SQ Sequence 1023 BP; 219 A; 272 C; 313 G; 219 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.:	5.28e-142	Length:	1023
Score:	1556.00	Matches:	318
Percent Similarity:	96.17%	Conservative:	8
Best Local Similarity:	93.81%	Mismatches:	13
Query Match:	90.36%	Indels:	3
DB:	8	Gaps:	0

US-10-088-079-2 (1-339) x ACA49224 (1-1023)

QY 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20

Db 1 ATGAACCAAGTAATGCGTCAATGACAGTCATCGTGCGGCTCGTACGGCACCT-CTC 59

QY 21 AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis 40

Db 60 GCCATCACTCTGGCGAGAAACGGCCACCAGGTTGCTGTGGGGCCACGCCAAACAT 119

QY 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProPhePro 60

Db 120 ATCGCGACCTGGAGCAGCATCGTGCAACGTGCGGTTCTTCCCGATGTGCTTTCCC 179

QY 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80

Db 180 GATACGTTACACTGGAAAGCGACTTAGCAACCGCGCTGGCGGCCAGTCGTAACATTCTG 239

QY 81 ValValValProSerHisValPheGlyGluValValLeuArgGlnIleLysProLeuMetArg 100

Db 240 GTGGTGGTGCCAAGCCCATGTTTTTCAGCGACGTGCTGCGGAGATTAAACCGCTGATCGGT 299

QY 101 ProAspAlaArgLeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120

Db 300 CCGGATGCGGCTCTGGTATGGGCGACCAAAAGGCTGGAAGCGGAAACGGGGCGCTGTTG 359

QY 121 GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140

Db 360 CAGGATGTCGCTCGCGAAGCGTTAGGCGATCAAAATCCCGCTGGCGGTGATTCTGGCCCG 419

QY 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160

Db 420 ACGTTCGCTAAAGAATTGGCGGCGGGTTTGGG-ACGGCAATCTCTCTGGCCTCAACCGAT 478

QY 161 GlnThrPheAlaAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180

Db 479 GAGACCTTTGGCGACGATCTCCAGCAACTGTTGCACCTGCGGAAAAAGTTTTCGCGTCTAT 538

QY 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle 200

Db 539 ATCAATGCGGATTTTATCGGCGTGACGTTGGCGGCGCGGTGAAAAACGTGATTGCGATT 598

QY 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220

Db 599 GGCGCGGGGATGCTGACGGCATCGGCTTCGGCGCGAAACGCCCGCACGCGCTAATCACG 658

QY 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240

Db 659 CGTGGAAGTACCGGAAATGTGCGGCTTGGCGCACG-CTTGGCGCGCATCCCGCCACCTTT 717

QY 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260

Db 718 ATGGGGATGGCGGGTTTAGGCGATCTGGTGCTGACCTGTACCGACAACAGTCGCGCAAC 777

QY 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280

Db 778 CGTCGTTTTGGCATGATGCTTGGCCAGGCGATGGACGTTAAAGGCGCGCAGGATAAGATT 837

QY 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300

Db 838 GGCCAGGTGTCGAAGGCTATCGCAATACGAAAGAAAGTTCTGTGAATTGGCGCACCGTTT 897

QY 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320

Db 898 GGTGTTGAAATGCCAATAACCGAGGAAATTTATCAAGTTTTTGTATTGCGGAAAAAACGCG 957

QY 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSerHis 339

Db 958 CGCGAGGCAGCATTAAACGTTATTAGTTCGCGCCCGCAAGGAAGAGCTGAGTCGCCAC 1014

RESULT 10

ACA54019

ID ACA54019 standard; DNA; 1020 BP.

XX

AC ACA54019;

XX

DT 19-JUN-2003 (first entry)

XX

DE Prokaryotic essential gene #35676.

XX

KW Antisense; ds; prokaryotic essential gene; cell proliferation;

KW drug design; gene.

XX

OS Yersinia pestis.

XX

PN WO200277183-A2.

XX

PD 03-OCT-2002.

XX

PF 21-MAR-2002; 2002WO-US009107.
XX
PR 21-MAR-2001; 2001US-00815242.
PR 06-SEP-2001; 2001US-00948993.
PR 25-OCT-2001; 2001US-0342923P.
PR 08-FEB-2002; 2002US-00072851.
PR 06-MAR-2002; 2002US-0362699P.
XX
PA (ELIT-) ELITRA PHARM INC.
XX
PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GU, Yamamoto R, Forsyth RA, Xu HH;
XX
DR WPI; 2003-029926/02.
DR P-PSDB; ABU50149.
XX
PT New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.
XX
XX Claim 14; SEQ ID NO 41889; 1766pp; English.
PS
CC The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
CC prokaryotic essential genes. Note: The sequence data for this patent did
CC not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 1020 BP; 238 A; 238 C; 288 G; 256 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 3.69e-132 Length: 1020
Score: 1455.00 Matches: 283
Percent Similarity: 91.37% Conservative: 24
Best Local Similarity: 84.23% Mismatches: 29
Query Match: 84.49% Indels: 0
DB: 8 Gaps: 0

US-10-088-079-2 (1-339) x ACA54019 (1-1020)

QY 1 MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu 20
Db 1 ATGAACACCAACCCTGCTTCAATGGCTGTTATCGGTGCGGATCTTACGGCACCGCATTA 60
QY 21 AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis 40
Db 61 GCTATCACACTGGCGGTAATGGCCATCAAGTCGTGTTATGGGGCCATGACCTAAACAT 120

QY 41 IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProPhePro 60
Db 121 ATTCAACAGCTGCAACAAGACCGCTGTAAACCGCGCTTTCCTACCTGATGCTGCTTCCCC 180
QY 61 AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu 80
Db 181 GATACGTTGCGATTGGAAACCGACTTAGCATGTGCGTTGGCTGCCAGCCGATGTGTG 240
QY 81 ValValValProSerHisValPheGlyGluValLeuAurGlnIleLysProLeuMetArg 100
Db 241 GTCGTCGTGCCAGCCATGTCTTTGGTGTGCTGTTTACATCAGTTGAAGCCTCATCTACGT 300
QY 101 ProAspAlaArgLeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu 120
Db 301 AAAGATGCACGTATCGTCTGGGCAACCAAGGGCTAGAAGCTGAACCGCGCTCTGCTA 360
QY 121 GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro 140
Db 361 CAGGATGTGGCCCGCGAAGTCTTGGGCGAGGCTATCCCGCTTCCGCTGATTTCTGGTCCA 420
QY 141 ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp 160
Db 421 ACGTTTGCCAAAGATTTGGCCGCGGTTTGCTTACGGCGATTCGTTGGCATCGACCGAT 480
QY 161 GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr 180
Db 481 GTGCAATTTAGCGAAGATCTGCAACAGTTATTGCACTGTGGAAGAGCTTTCGAGTTTAC 540
QY 181 SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle 200
Db 541 AGTAATCCTGATTTTATCGGGGTACAGCTTGGTGGCGCAGTGAAACACGTGATTGCCATC 600
QY 201 GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr 220
Db 601 GGTGCAGGTATGTCGATGGCATCGGTTTGGTGGCAATGCCGTACCGCTCTAATAACC 660
QY 221 ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe 240
Db 661 CGCGGGTTAGCGGAGATGACGCGCTTAGGGACGCGATTAGGTGCCGATCCTTCCACCTTT 720
QY 241 MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn 260
Db 721 ATGGGCATGGCAGGGTTAGCGGATTTGGTGTCTAACCTGCACAGATAACCAATCCCGTAAC 780
QY 261 ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle 280
Db 781 CGCCGATTTGGCATTTATGCTGGGTACAGGGGTTGGGGGTGAAGGAGGCGCAGGACAACATT 840
QY 281 GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe 300
Db 841 GGTCAAGTGGTAGAAGGTTACCGTAATACCAAGGAAGTTCTGGCATTAGCACAGCGTCAT 900
QY 301 GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla 320
Db 901 GGCGTCGAAATGCCAATAACTGAACAAATTTATCAAGTGTGTATTGTTCATAAGAATGCT 960
QY 321 ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGluArg 336
Db 961 CGTGAGGCGGCTCTGACGTTGTTGGGGCGGACCAAAAAAGATGAAAAA 1008
RESULT 11
ADF03046
ID ADF03046 standard; DNA; 1023 BP.
XX
AC ADF03046;
XX
DT 12-FEB-2004 (first entry)
XX
DE Bacterial polynucleotide #3331.
XX
KW Proteus mirabilis infection; bacterial infection; antibacterial;
KW immunostimulant; gene; ds.

XX OS Proteus mirabilis.
XX PN US6605709-B1.
XX XX
PD 12-AUG-2003.
XX XX
PF 05-APR-2000; 2000US-00543681.
XX XX
PR 09-APR-1999; 99US-0128706P.
XX XX
PA (GENO-) GENOME THERAPEUTICS CORP.
XX XX
PI Breton GL;
XX XX
DR WPI; 2003-895291/82.
DR P-PSDB; ADF07218.
XX XX
PT New Proteus mirabilis polypeptides and polynucleotides, useful as
PT reagents for diagnosis of bacterial disease, as components of
PT antibacterial vaccines, as targets for antibacterial drugs, or as
PT biocontrol agents for plants.
XX XX
PS Disclosure; SEQ ID NO 3331; 870pp; English.
XX XX
CC The invention relates to new Proteus mirabilis polypeptides and
CC polynucleotides. The invention also relates to antibodies against the
CC polypeptides, methods for producing the polypeptides, a method of
CC generating vaccines for immunising an individual against P. mirabilis, a
CC method for evaluating a compound for the ability to bind a P. mirabilis
CC polypeptide and a method for screening test compounds for anti-bacterial
CC activity. The polypeptides and polynucleotides are useful as molecular
CC targets for diagnosing, preventing and treating pathological conditions
CC resulting from bacterial infection, as reagents for diagnosis of
CC bacterial diseases, as components of antibacterial vaccines, as targets
CC for antibacterial drugs or as bio-control agents for plants. This
CC sequence represents a Proteus mirabilis polynucleotide of the invention.
XX XX
SQ Sequence 1023 BP; 266 A; 206 C; 264 G; 287 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 9.56e-129 Length: 1023
Score: 1420.00 Matches: 275
Percent Similarity: 90.72% Conservative: 28
Best Local Similarity: 82.34% Mismatches: 31
Query Match: 82.46% Indels: 0
DB: 10 Gaps: 0

US-10-088-079-2 (1-339) x ADF03046 (1-1023)

QY 5 AsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeuAlaIleThrLeu 24
Db 13 AACGCTTCTATGACAGTATCGGTGCCGTTCAACGGCACCGCTTTAGCGATTACCTTA 72
QY 25 AlaArgAsnGlyHisGluValValLeuTyrGlyHisAspProGluHisIleAlaThrLeu 44
Db 73 GCGGTAATGGGCACGATGTTGTCTGTGGGGGATGATCCCAAGCACGTTGGGCATTA 132
QY 45 GluArgAspArgCysAsnAlaAlaPheLeuProAspValPropheProAspThrLeuHis 64
Db 133 GAACAAGCGCGCTGTATCAAGCCTTCTGCCTGATGTTCCCTTCCTGATAGTTTATAT 192
QY 65 LeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeuValValPro 84
Db 193 ATGGAAGCTTCTTTGCAAAAAGCGATTGAAGCGAGCGGTAATATTTCTTGTCGATCCCA 252
QY 85 SerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArgProAspAlaArg 104
Db 253 AGCCATGTGTTGGTGAAGTACTGCAACAAATCAAAACCCCTTTTACGTCAGGATGCGCGT 312
QY 105 LeuValTyrAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeuGlnAspValAla 124
Db 313 GTTGTGTTGGCGCACAAAAGGTCTTTGAAGCACATACACTGGTTCGCTTATTACAAGATGTTGCC 372

QY 125 ArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyProThrPheAlaLys 144
Db 373 CGTGAAGTATTAGGTAATGAATCCCGCTCGCAGTATTATCTGGCCCTACTTTTGTGTA 432
QY 145 GluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAspGlnThrPheAla 164
Db 433 GAGCTTGCCGCGGTTTGCCACCGCGATTGCGAGTGGCTCGACGGATAATCTCTTTTA 492
QY 165 AspAspLeuGlnLeuLeuHisCysGlyLysSerPheArgValTyrSerAsnProAsp 184
Db 493 GAGCAGTTACACAGCTATTTTCATTGTGTGTAAGTTTCCGAGTCTATAAAAACCTGAT 552
QY 185 PheIleGlyValGlnLeuGlyAlaValLysAsnValIleAlaIleGlyAlaGlyMet 204
Db 553 TTTATCGGTGTGCAACTGGTGGCGCTGTAAAAACGTGATTGCTATTGTCGGGTATG 612
QY 205 SerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThrArgGlyLeuAla 224
Db 613 TCTGATGGTATGGGATTGTCGCTAATGCGCGTACCGCACTGATTACCGAGGTCTAGCC 672
QY 225 GluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPheMetGlyMetAla 244
Db 673 GAAATGAGCCGTTAGTAAAGCCTTAGGTGCAGATGCGCACTTTTATGGGCATGGCT 732
QY 245 GlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsnArgArgPheGly 264
Db 733 GGTGTTGGGTGATTAGTTTAACTGTACTGACAACCAATCACGTAATCGTCGCTTCGGT 792
QY 265 MetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnLysIleGlyGlnValVal 284
Db 793 ATGATGCTAGGTCAAGTTTAGATTGTGATACAGCCCAAGAGAAAATTGSCCAGGTAGTC 852
QY 285 GluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPheGlyValGluMet 304
Db 853 GAAGGTTATCGTAACACCAAGAAGTTCCGCGCATTAGCCGAACAGGTGGTGTAGAAATG 912
QY 305 ProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAlaArgGluAlaAla 324
Db 913 CCAATCACCGAACAGATCTACCAATTTTATATCAACATAAAGATGTAAAAAGAGGCTGCA 972
QY 325 LeuThrLeuLeuGlyArgAlaArgLysAspGluArgSerSer 338
Db 973 TTGGCTTTATTAGGCGGAGCAACCAAGATGAGATAGACAGC 1014
RESULT 12
ACA44463
ID ACA44463 standard; DNA; 1011 BP.
XX
AC ACA44463;
XX
DT 19-JUN-2003 (first entry)
XX
DE Prokaryotic essential gene #26120.
XX
KW Antisense; ds; prokaryotic essential gene; cell proliferation;
KW drug design; gene.
XX
OS Proteus sp.
XX
PN WO200277183-A2.
XX
PD 03-OCT-2002.
PF 21-MAR-2002; 2002WO-US009107.
XX
PR 21-MAR-2001; 2001US-00815242.
PR 06-SEP-2001; 2001US-00948993.
PR 25-OCT-2001; 2001US-0342923P.
PR 08-FEB-2002; 2002US-00072851.
XX 06-MAR-2002; 2002US-0362699P.
PA (ELIT-) ELITRA PHARM INC.

XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
XX
DR WPI; 2003-029926/02.
DR P-PSDB; ABU40593.
XX
PT New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.
XX
PS Claim 14; SEQ ID NO 32333; 1766pp; English.
XX
CC The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
CC prokaryotic essential genes. Note: The sequence data for this patent did
CC not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 1011 BP; 260 A; 206 C; 261 G; 284 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 7.1e-128 Length: 1011
Score: 1411.00 Matches: 274
Percent Similarity: 90.12% Conservative: 27
Best Local Similarity: 82.04% Mismatches: 33
Query Match: 81.94% Indels: 0
DB: 8 Gaps:

US-10-088-079-2 (1-339) x ACA44463 (1-1011)

QY 5 AsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeuAlaIleThrLeu 24
Db 4 AACGCTCTATGACAGTTATCGGTGCCGGTTTCATACGGCACCGCTTAGCGATTACCTTA 63

QY 25 AlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHisIleAlaThrLeu 44
Db 64 GCGCGTAATGGGCACGATGTTGTGCTGTGGGGCATGATCCCAAGCACGTTGCCGGCATTA 123

QY 45 GluArgAspArgCysAsnAlaAlaPheLeuProAspValProPheProAspThrLeuHis 64
Db 124 GAACAAGCGCGCTGTAATCAAGCCTTCTGCCCGATGTTTCCTTCCTGATAGTTTATAT 183

QY 65 LeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeuValValPro 84
Db 184 ATGGAAGCTTCTTTGCAAAAAGCGGATGAAGCGAGCGCGTAATATCTTGTGGTGATCCCA 243

QY 85 SerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArgProAspAlaArg 104
Db 244 AGCCATGTGTTTGGTGAAGTACTGCAACAAATCAAACCCCTTTTACGTACAGGATGCGCGT 303

QY 105 LeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuGlnAspValAla 124
Db 304 GTTGTTTGGGCGACAAAAGGTCTTTGAAGCACATACTGTCGCTTATTACAAGATGTTGCC 363

QY 125 ArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyProThrPheAlaLys 144
Db 364 CGTGAAGTATTAGGTAATGAAATCCCGCTCGCAGTATTATCTGGCCCTACTCTTTGCTAAA 423

QY 145 GluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAspGlnThrPheAla 164
Db 424 GAGCTTGCAGCGGTTTACCCACCGGATTCAGTGGCTCGACGGATAATCTCTTTTA 483

QY 165 AspAspLeuGlnGlnLeuHisCysGlyLysSerPheArgValTyrSerAsnProAsp 184
Db 484 GAACAGTTACAACAGCTATTTTCATTGTGTGTAAGCTCCGAGTCTATAAAACCCCTGAT 543

QY 185 PheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIleGlyAlaGlyMet 204
Db 544 TTTATCGGTGTGCAACTGGGGGCGCTGTTTAAAAACGTGATTGCTATTGGTGGGGTATG 603

QY 205 SerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThrArgGlyLeuAla 224
Db 604 TCTGATGGTATGGGATTTGGCGCTAATGCGGCTACTGCACCTATTACCCGTGGTCTGGCC 663

QY 225 GluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPheMetGlyMetAla 244
Db 664 GAAATGAGCCGCTTAGGTAAAGCCTTAGGTGCAGATGCGGCAACTTTTATGGGCATGGCT 723

QY 245 GlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsnArgArgPheGly 264
Db 724 GGTTTGGGTGATTTAGTTTAACTGTACTGACAAACCAATCAGTAATCGTCGCTTCGGT 783

QY 265 MetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIleGlyGlnValVal 284
Db 784 ATGATGCTAGGTCAAGGTTTGGTGTGTGATACAGCCCAAGAGAAAATTGGCCAGGTAGTC 843

QY 285 GluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPheGlyValGluMet 304
Db 844 GAAGGTTATCGTTAACCAACCAAGAAAGTTTCGCGCATTAGCCGAACAGGTGGGTGTAGAAATG 903

QY 305 ProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAlaArgGluAlaAla 324
Db 904 CCAATCACCGAACAGATCTACCAATTTTATATCAACATAAAGATGTAAAAGAGSGCTGCA 963

QY 325 LeuThrLeuLeuGlyArgAlaAlaArgLysAspGluArgSerSer 338
Db 964 TTGGCTTTATTAGGGCGAGCAACCAAGATGAGATAGACAGC 1005

RESULT 13
ACF70491
ID ACF70491 standard; DNA; 1023 BP.
XX
AC ACF70491;
XX
DT 20-NOV-2003 (first entry)
XX
DE Photorhabdus luminescens nucleotide sequence #8958.
XX
KW Antibacterial; fungicide; insecticide; polymorphism; genetic analysis;
KW detection; food; gene expression; plant; animal; microorganism; toxin;
KW antibiotic; biopesticide; virulence factor; disease model; plague;
KW whooping cough; gene; ds.
XX
OS Photorhabdus luminescens.
XX
PN WO200294867-A2.
XX
PD 28-NOV-2002.
XX

XX Duchaud E, Taourit S, Glaser P, Frangeul L, Kunst F, Danchin A;
PI Buchrieser C;
XX
DR WPI; 2003-148459/14.
XX
XX Genomic sequence of Photorhabdus luminescens and encoded polypeptides,
PT useful e.g. as therapeutic antimicrobials and agricultural pesticides.
XX
XX Claim 1; SEQ ID NO 27; 1205pp; French.
PS
XX
CC The invention relates to the isolation of genes and their encoded
CC proteins from Photorhabdus luminescens. The isolated sequences are
CC sources of probes and primers for detecting the genome of P. luminescens
CC and related species; to study polymorphisms; for gene analysis and for
CC detection/amplification of the genes. Antibodies (Ab) raised against the
CC polypeptides encoded by the genes are used for detection/identification
CC of P. luminescens, e.g. in foods. The genes, proteins, Ab and cells that
CC carry a gene-containing vector are used to select compounds that
CC modulate, regulate, induce or inhibit expression of the genes in plants,
CC animals or microorganisms other than P. luminescens and are able to alter
CC response or sensitivity to toxins and antibiotics produced by P.
CC luminescens. Cells transformed to express the genes are useful for
CC recombinant production of the proteins, particularly toxins and
CC antibacterials useful as insecticides, bactericides and fungicides. The
CC genes, proteins, vectors containing the genes and Ab are also useful
CC therapeutically (to treat microbial infection by bacteria or fungi that
CC are sensitive to P. luminescens-encoded toxins or antibiotics) and as
CC biopesticides. Other uses of the genes and the proteins are as virulence
CC factors and for identifying targets of human diseases for which P.
CC luminescens is a model (particularly plague and whooping cough). This
CC sequence represents one of the isolated P. luminescens genes
XX
SQ Sequence 69727 BP; 20213 A; 13239 C; 14632 G; 21638 T; 0 U; 5 Other;

Alignment Scores:
Pred. No.: 9.67e-124 Length: 69727
Score: 1393.50 Matches: 270
Percent Similarity: 89.55% Conservative: 30
Best Local Similarity: 80.60% Mismatches: 34
Query Match: 80.92% Indels: 1
DB: 10 Gaps: 1

US-10-088-079-2 (1-339) x ACF65374 (1-69727)

Qy	1	MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu	20
Db	54175	ATGAATAGT---ACTGTTTCTATGACAGTGAATTGGTGGCGCTCATACGGCACCTCATTA	54231
Qy	21	AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis	40
Db	54232	GCCATTACGCTGGCTCGTAATGTCATAATGTTGTACTTTGGGGGCATAATCCAGAGCAT	54291
Qy	41	IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProPhePro	60
Db	54292	GTGGGGCATTGCAACGGGTGCTTGTAAATCAAAAATTTCTGCCGGATGTTTCCTTCCT	54351
Qy	61	AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu	80
Db	54352	GATAGTTTATTGCTTGAAACGGACCTAATAAAGCACTAACAGCGAGCCCGGATATCTT	54411
Qy	81	ValValValProSerHisValPheGlyGluValLeuArgGlnIleLysProLeuMetArg	100
Db	54412	GTTGGGTACCTAGCCATGTTGTTGGTGAAGTGTAAAGCAGAGATAAAACCACATTACGG	54471
Qy	101	ProAspAlaArgLeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu	120
Db	54472	CCTGATTACGTCATCGTATGGCAACTAAAGGCTTGAAGCGGATACCGGTGCGTTATTG	54531
Qy	121	GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro	140
Db	54532	CAGGATGTGGCCCGTGAGATATTAGGCAATGAAATACCGTAGCGGTGCTCTCTGGGCCA	54591

Qy	141	ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp	160
Db	54592	ACATTTGCTAAAGAGTTAGCGGCTGGTTTGCCCTACCGGATTGCTATTTCCGCGACGGAA	54651
Qy	161	GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr	180
Db	54652	TCTGCTTTTGGCGATGACTTCAACAATATTATCCACTGTGGCAAAAGTTTCCGGGTTTAT	54711
Qy	181	SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle	200
Db	54712	AAAAATCCTGATTTTATTGGTGTTCAACTCGGTGGTCCGTAATAAACCGTGATCGCCATT	54771
Qy	201	GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr	220
Db	54772	GGCGCGGGAATATCTGATGGCATGGGATTTGGTGTCTTAATGCTCGTACCGCATTTACT	54831
Qy	221	ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe	240
Db	54832	CGTGGATTGGCGGAAATGAGTCGCCTTGGTGCAGCGCTTGGTGTGATCCTTCTACCTTT	54891
Qy	241	MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn	260
Db	54892	ATGGGCATGGCGGATTTGGCGATTTGGTCTTAACTTGTTACTGTATAACCAATCACGTAAC	54951
Qy	261	ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle	280
Db	54952	CGTCGTTTGGCATGATGCTGGGCGAGGAATCAGTGTTTGAAGAAGCGCAGTATCAGATT	55011
Qy	281	GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe	300
Db	55012	GGGCAGGTTGTTGAAGTTATCGCAATACCAAGAGTAGCTGTCATTGGCTAATCGCGCC	55071
Qy	301	GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla	320
Db	55072	AATGTAGAAATGCCGATTGCAGAACAAATCTACCAGATACTCTATTGCAATAAAATGTG	55131
Qy	321	ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGlu	335
Db	55132	ATAGAAGCTGCTCAGGCATTATTAGGAAGAGCCAGAAAGGATGAG	55176

RESULT 15

ACF67367_35

Continuation (36 of 57) of ACF67367 from base 3500001 (Photorhabdus luminescens nucleoti
WP Sequence split into 57 fragments LOCUS ACF67367 Accession Acf67367

WP	Fragment Name	Begin	End
WP	ACF67367_00	1	110000
WP	ACF67367_01	100001	210000
WP	ACF67367_02	200001	310000
WP	ACF67367_03	300001	410000
WP	ACF67367_04	400001	510000
WP	ACF67367_05	500001	610000
WP	ACF67367_06	600001	710000
WP	ACF67367_07	700001	810000
WP	ACF67367_08	800001	910000
WP	ACF67367_09	900001	1010000
WP	ACF67367_10	1000001	1110000
WP	ACF67367_11	1100001	1210000
WP	ACF67367_12	1200001	1310000
WP	ACF67367_13	1300001	1410000
WP	ACF67367_14	1400001	1510000
WP	ACF67367_15	1500001	1610000
WP	ACF67367_16	1600001	1710000
WP	ACF67367_17	1700001	1810000
WP	ACF67367_18	1800001	1910000
WP	ACF67367_19	1900001	2010000
WP	ACF67367_20	2000001	2110000
WP	ACF67367_21	2100001	2210000
WP	ACF67367_22	2200001	2310000
WP	ACF67367_23	2300001	2410000
WP	ACF67367_24	2400001	2510000
WP	ACF67367_25	2500001	2610000
WP	ACF67367_26	2600001	2710000
WP	ACF67367_27	2700001	2810000

WP	ACF67367_28	2800001	2910000
WP	ACF67367_29	2900001	3010000
WP	ACF67367_30	3000001	3110000
WP	ACF67367_31	3100001	3210000
WP	ACF67367_32	3200001	3310000
WP	ACF67367_33	3300001	3410000
WP	ACF67367_34	3400001	3510000
WP	ACF67367_35	3500001	3610000
WP	ACF67367_36	3600001	3710000
WP	ACF67367_37	3700001	3810000
WP	ACF67367_38	3800001	3910000
WP	ACF67367_39	3900001	4010000
WP	ACF67367_40	4000001	4110000
WP	ACF67367_41	4100001	4210000
WP	ACF67367_42	4200001	4310000
WP	ACF67367_43	4300001	4410000
WP	ACF67367_44	4400001	4510000
WP	ACF67367_45	4500001	4610000
WP	ACF67367_46	4600001	4710000
WP	ACF67367_47	4700001	4810000
WP	ACF67367_48	4800001	4910000
WP	ACF67367_49	4900001	5010000
WP	ACF67367_50	5000001	5110000
WP	ACF67367_51	5100001	5210000
WP	ACF67367_52	5200001	5310000
WP	ACF67367_53	5300001	5410000
WP	ACF67367_54	5400001	5510000
WP	ACF67367_55	5500001	5610000
WP	ACF67367_56	5600001	5648894

Alignment Scores:

Pred. No.:	1.77e-123	Length:	110000
Score:	1393.50	Matches:	270
Percent Similarity:	89.55%	Conservative:	30
Best Local Similarity:	80.60%	Mismatches:	34
Query Match:	80.92%	Indels:	1
DB:	10	Gaps:	1

US-10-088-079-2 (1-339) x ACF67367_35 (1-110000)

QY	1	MetAsnGlnArgAsnAlaSerMetThrValIleGlyAlaGlySerTyrGlyThrAlaLeu	20
DB	79826	ATGAATAGT---ACTGTTTCTATGACAGTGATTGGTCCGGCTCATACGGCACCTCATT	79882
QY	21	AlaIleThrLeuAlaArgAsnGlyHisGluValValLeuTrpGlyHisAspProGluHis	40
DB	79883	GCCATTACGCTGGCTCGTAATGGTGCATAATGTTGTAATCAAAAATTTCTGCCGGATGTTTCCTT	79942
QY	41	IleAlaThrLeuGluArgAspArgCysAsnAlaAlaPheLeuProAspValProphePro	60
DB	79943	GTTGGGGCATTGCAACGGGTGCGTTGTAATCAAAAATTTCTGCCGGATGTTTCCTT	80002
QY	61	AspThrLeuHisLeuGluSerAspLeuAlaThrAlaLeuAlaAlaSerArgAsnIleLeu	80
DB	80003	GATAGTTTATTGCTTGAAACGGACCTAATAAAGCACTAACAGCGAGCCCGCATATTCTT	80062
QY	81	ValValValProSerHisValPheGlyGluValValLeuArgGlnIleLysProLeuMetArg	100
DB	80063	GTTGTGGTACCTAGCCATGTGTTTGGTGAAGTGTAAAGCAGATAAAACCATTTACGG	80122
QY	101	ProAspAlaArgLeuValTrpAlaThrLysGlyLeuGluAlaGluThrGlyArgLeuLeu	120
DB	80123	CCTGATTCACGTATCGTATGGGCAACTAAAGGCTTGGAAAGCGGATACCGGTCGGTTATTG	80182
QY	121	GlnAspValAlaArgGluAlaLeuGlyAspGlnIleProLeuAlaValIleSerGlyPro	140
DB	80183	CAGGATGTGGCCCGTGAGATATTAGGCAATGAATACCGTAGCGGTGCTCTCTGGGCCA	80242
QY	141	ThrPheAlaLysGluLeuAlaAlaGlyLeuProThrAlaIleSerLeuAlaSerThrAsp	160
DB	80243	ACATTTGCTAAAGAGTTAGCGGCTGGTTTGCTACCGCATTGCTATTTCCGCGACGGAA	80302
QY	161	GlnThrPheAlaAspAspLeuGlnGlnLeuLeuHisCysGlyLysSerPheArgValTyr	180

Db	80303	TCTGCTTTTGGCGATGGACTTCAACAATATTATTCACACTGTGGCAAAAGTTTCCGGGTTTAT	80362
QY	181	SerAsnProAspPheIleGlyValGlnLeuGlyGlyAlaValLysAsnValIleAlaIle	200
Db	80363	AAAAATCCTGATTTATTGGTGTTCAACTCGGTGGTCCCGTAAAAAACGTGATCGCCATT	80422
QY	201	GlyAlaGlyMetSerAspGlyIleGlyPheGlyAlaAsnAlaArgThrAlaLeuIleThr	220
Db	80423	GGCGCGGAATATCTGATGGCATGGGATTTGGTGCTAATGCTCGTACCGCATTTGATTACT	80482
QY	221	ArgGlyLeuAlaGluMetSerArgLeuGlyAlaAlaLeuGlyAlaAspProAlaThrPhe	240
Db	80483	CGTGGATTGGCGAAATGAGTCGCCCTTGGTGACGCGCTTGGTGCTGATCTCTACCTTT	80542
QY	241	MetGlyMetAlaGlyLeuGlyAspLeuValLeuThrCysThrGluAsnGlnSerArgAsn	260
Db	80543	ATGGGCATGGCGGATTTGGCGGATTTGGTCTTAACTTGTAAGAACGCGCAGTATCAGATT	80602
QY	261	ArgArgPheGlyMetMetLeuGlyGlnGlyMetAspValGlnSerAlaGlnGluLysIle	280
Db	80603	CGTCGTTTGGCATGATGCTGGGCGAGGAATCAGTGTGAAGAACGCGCAGTATCAGATT	80662
QY	281	GlyGlnValValGluGlyTyrArgAsnThrLysGluValArgGluLeuAlaHisArgPhe	300
Db	80663	GGGCAGGTTGTGAAGGTTATCGCAATACCAAGAAAGTACGTCGATTGGCTAATCGCGCC	80722
QY	301	GlyValGluMetProIleThrGluGluIleTyrGlnValLeuTyrCysGlyLysAsnAla	320
Db	80723	AATGTAGAAATGCCGATTGCAGAACAAATCTACCAGATACTCTATTGCAATAAAATGTG	80782
QY	321	ArgGluAlaAlaLeuThrLeuLeuGlyArgAlaArgLysAspGlu	335
Db	80783	ATAGAAAGCTGCTCAGGCATTATTAGGAAGAGCCAGAAAGGATGAG	80827

Search completed: April 27, 2005, 15:42:11
Job time : 772 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: April 27, 2005, 10:44:58 ; Search time 164 Seconds
(without alignments)
799.462 Million cell updates/sec

Title: US-10-088-079-2
Perfect score: 1722
Sequence: 1 MNQRNASMTVIGAGSYGTAL.....AREAAALTLGLRARKDERSH 339

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues

Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_16Dec04:*
1: geneseqp19808:*
2: geneseqp19908:*
3: geneseqp20008:*
4: geneseqp20018:*
5: geneseqp20028:*
6: geneseqp2003as:*
7: geneseqp2003bs:*
8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	1722	100.0	339	4	AAB62189	Aab62189 E. coli g
2	1719	99.8	339	2	AAW57330	Aaw57330 Glycerol-
3	1719	99.8	339	2	AAW60258	Aaw60258 Klebsiell
4	1719	99.8	339	2	AAY26172	Aay26172 Glycerol-
5	1719	99.8	339	4	AAU34796	Aau34796 E. coli c
6	1719	99.8	339	6	ABU28819	Abu28819 Protein e
7	1719	99.8	339	8	ADS45174	Ads45174 Bacterial
8	1640	95.2	339	6	ABU47465	Abu47465 Protein e
9	1635	94.9	339	6	ABU28096	Abu28096 Protein e
10	1621	94.1	339	6	ABU45354	Abu45354 Protein e
11	1587	92.2	339	6	ABU31968	Abu31968 Protein e
12	1587	92.2	345	7	ABO64223	AbO64223 Klebsiell
13	1455	84.5	339	6	ABU50149	Abu50149 Protein e
14	1420	82.5	340	7	ADF07218	Adf07218 Bacterial
15	1411	81.9	337	6	ABU40593	Abu40593 Protein e
16	1393.5	80.9	341	6	ABM68792	Abm68792 Photorhab
17	1373	79.7	330	8	ADN17703	Adn17703 Bacterial
18	1372	79.7	330	8	ADS42893	Ads42893 Bacterial
19	1242	72.1	344	6	ABU49620	Abu49620 Protein e
20	1192	69.2	337	6	ABU39371	Abu39371 Protein e
21	1150	66.8	335	4	AAU35472	Aau35472 Haemophil
22	1150	66.8	335	6	ABU30312	Abu30312 Protein e
23	806	46.8	329	6	ABU33190	Abu33190 Protein e
24	794.5	46.1	334	8	ADS26951	Ads26951 Bacterial
25	794.5	46.1	334	8	ADS27309	Ads27309 Bacterial

26	794.5	46.1	340	8	ADS26576	Ads26576 Bacterial
27	751.5	43.6	346	8	ADS28582	Ads28582 Bacterial
28	738.5	42.9	334	8	ADN26503	Adn26503 Bacterial
29	720.5	41.8	334	8	ADS27486	Ads27486 Bacterial
30	709.5	41.2	340	6	ABU18125	Abu18125 Protein e
31	703	40.8	332	6	ABU23655	Abu23655 Protein e
32	694	40.3	343	6	ABU25613	Abu25613 Protein e
33	692	40.2	355	6	ABU30030	Abu30030 Protein e
34	692	40.2	356	7	ADC94377	Adc94377 E. faeciu
35	681	39.5	353	7	ADC95426	Adc95426 E. faeciu
36	677.5	39.3	339	6	ADB08818	Adb08818 Alloiococ
37	677.5	39.3	342	6	ADB08820	Adb08820 Alloiococ
38	673	39.1	340	4	AAU35259	Aau35259 Enterococ
39	673	39.1	340	6	ABU14598	Abu14598 Protein e
40	672.5	39.1	338	5	ABB49213	Abb49213 Listeria
41	672.5	39.1	338	6	ABU32661	Abu32661 Protein e
42	672.5	39.1	342	4	AAU33428	Aau33428 Enterococ
43	666.5	38.7	327	6	ADB08816	Adb08816 Alloiococ
44	666.5	38.7	345	8	ADS44752	Ads44752 Bacterial
45	666	38.7	335	6	ABU17729	Abu17729 Protein e

ALIGNMENTS

RESULT 1

AAB62189 ID AAB62189 standard; protein; 339 AA.

XX AC AAB62189;

XX DT 11-JUN-2001 (first entry)

XX XX E. coli gpsA2FR protein.

XX DE Glycerol-3-phosphate dehydrogenase; G3PD; feedback inhibition; oil seed;

KW KW Genetic transformation; fatty acid; glycerolipid; osmotic stress; gpsA;

XX KW gpsA2FR; allele.

OS Escherichia coli.

XX FH Key Location/Qualifiers

FT Misc-difference 255

FT FT /label= D255E

XX FT /note= "wild-type Asp is replaced with Glu"

PN WO200121820-A1.

XX PD 29-MAR-2001.

XX PF 21-SEP-2000; 2000WO-CA001096.

XX PR 22-SEP-1999; 99US-0155133P.

XX PA (CANA) NAT RES COUNCIL CANADA.

XX PI Zou J, Wei Y, Periappuram C; Selvaraj G, Datla R;

XX DR WPI; 2001-257996/26.

XX DR N-PSDB; AAF57428.

XX PT Manipulating glycerol-3-phosphate metabolism of plant for enhancing

PT stress tolerance, altering fatty acid content in glycerolipids, by

PT expressing in plant feedback defective glycerol-3-phosphate dehydrogenase

XX gene.

XX PS Claim 15; Fig 1; 39pp; English.

XX CC The invention provides a method for genetically transforming a plant so

CC that it expresses a heterologous glycerol-3-phosphate dehydrogenase

CC (G3PD) that is less sensitive to feedback inhibition than wild-type G3PD.

CC The method involves providing a vector comprising a DNA sequence encoding

CC G3PD that is less sensitive to feedback inhibition than wild-type G3PD

CC and transforming the plant with the vector. The method is useful for
CC expressing a heterologous G3PD less sensitive to feedback inhibition than
CC wild-type G3PD in an oil seed bearing plant, such as *Arabidopsis thaliana*
CC or *Brassica*. The vectors are useful for producing a genetically altered
CC plant having altered fatty acid content in its glycerolipids, especially
CC elevated levels of C16 fatty acids and increased osmotic stress tolerance
CC relative to the wild type. The present sequence represents the *E. coli*
CC *gpsA2FR* protein. The gene *gpsA2FR* is an allele of the *E. coli* *gpsA* gene,
CC and encodes an altered version of the GPDH protein defective in feedback
CC inhibition. This *gpsA2FR* gene can be used in the vectors and method of
CC the invention
XX
SQ Sequence 339 AA;

Query Match 100.0%; Score 1722; DB 4; Length 339;
Best Local Similarity 100.0%; Pred. No. 7.2e-159;
Matches 339; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEHIATLERDRCNAAFLPDVPFP 60
|||
Db 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEHIATLERDRCNAAFLPDVPFP 60

QY 61 DTLHLESDLATALAASRNILVVVPSHVGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
|||
Db 61 DTLHLESDLATALAASRNILVVVPSHVGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120

QY 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFVY 180
|||
Db 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFVY 180

QY 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATF 240
|||
Db 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATF 240

QY 241 MGMAGLDLVLCTENQSRNRRFGMMLGQGM DVQSAQEKIGQVVEGYRNTKEVRELAHRF 300
|||
Db 241 MGMAGLDLVLCTENQSRNRRFGMMLGQGM DVQSAQEKIGQVVEGYRNTKEVRELAHRF 300

QY 301 GVEMPTIEEIIQVLYCGKNAREAAALTLGRARKDERSH 339
|||
Db 301 GVEMPTIEEIIQVLYCGKNAREAAALTLGRARKDERSH 339

RESULT 2
AAW57330
ID AAW57330 standard; protein; 339 AA.
XX
AC AAW57330;
XX
DT 14-SEP-1998 (first entry)
XX
DE Glycerol-3-phosphate dehydrogenase *gpsA*.
XX
KW Glycerol-3-phosphate dehydrogenase; G3PDH; *gpsA*; yeast.
XX
OS *Saccharomyces* sp.
XX
PN WO9821340-A1.
XX
PD 22-MAY-1998.
XX
PF 10-NOV-1997; 97WO-US020293.
XX
PR 13-NOV-1996; 96US-0030602P.
XX
PA (DUPO) DU PONT DE NEMOURS & CO E I.
PA (GEMV) GENENCOR INT INC.
XX
PI Bulthuis BA, Gatenby AA, Haynie SL, Hsu AK, Lareau RD;
XX
DR WPI; 1998-297943/26.
XX
PT Fermentative production of glycerol using recombinant host - containing

PT genes for glycerol-3-phosphate dehydrogenase and-or glycerol-3-
PT phosphatase.
XX
PS Claim 9; Page 36-37; 57pp; English.
XX
CC This claimed *Saccharomyces* polypeptide comprises a glycerol-3-phosphate
CC dehydrogenase (G3PDH) that catalyses the conversion of dihydroxyacetone
CC phosphate to glycerol-3-phosphate. It is encoded by the *gpsA* gene. The
CC invention provides recombinant organisms that express G3PDH and/or
CC glycerol-3-phosphatase (G3P) (see also AAW57324-32) useful for the
CC production of glycerol from a variety of C-sources. A host cell is
CC preferably transformed with a cassette containing either a G3PDH gene
CC and/or a G3P gene and then cultured in the presence of a mono-, oligo-,
CC polysaccharide or 1C-substrate. The glycerol obtained is used in
CC cosmetics, liquid soaps, pharmaceuticals, lubricants and antifreezes; its
CC esters are used in the oil and fat industries. The method produces
CC glycerol rapidly and inexpensively without generation of polluting by-
CC products
XX
SQ Sequence 339 AA;

Query Match 99.8%; Score 1719; DB 2; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.4e-158;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEHIATLERDRCNAAFLPDVPFP 60
|||
Db 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEHIATLERDRCNAAFLPDVPFP 60

QY 61 DTLHLESDLATALAASRNILVVVPSHVGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
|||
Db 61 DTLHLESDLATALAASRNILVVVPSHVGEVLRQIKPLMRPDARLVWATKGLEAETGRLL 120

QY 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFVY 180
|||
Db 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLLHCGKSFVY 180

QY 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATF 240
|||
Db 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATF 240

QY 241 MGMAGLDLVLCTENQSRNRRFGMMLGQGM DVQSAQEKIGQVVEGYRNTKEVRELAHRF 300
|||
Db 241 MGMAGLDLVLCTENQSRNRRFGMMLGQGM DVQSAQEKIGQVVEGYRNTKEVRELAHRF 300

QY 301 GVEMPTIEEIIQVLYCGKNAREAAALTLGRARKDERSH 339
|||
Db 301 GVEMPTIEEIIQVLYCGKNAREAAALTLGRARKDERSH 339

RESULT 3
AAW60258
ID AAW60258 standard; protein; 339 AA.
XX
AC AAW60258;
XX
DT 28-SEP-1998 (first entry)
XX
DE *Klebsiella pneumoniae* glycerol-3-phosphate dehydrogenase.
XX
KW glycerol-3-phosphate dehydrogenase; production; 1,3-propanediol;
KW recombinant.
XX
OS *Klebsiella pneumoniae*.
XX
PN WO9821341-A2.
XX
PD 22-MAY-1998.
XX
PF 13-NOV-1997; 97WO-US020873.
XX
PR 13-NOV-1996; 96US-0030601P.
XX

PA (GEMV) GENENCOR INT INC.
XX
PI Dunn-Coleman NS, Diaz-Torres M, Chase MW, Trimbur D;
XX
XX WPI; 1998-297944/26.
DR
XX
XX New method for increasing production of 1,3-propane:diol - comprises
PT fermentation of inexpensive carbon sources by microorganism expressing
PT dehydratase, used, e.g. to prolong half-life of enzyme.
XX
PS Disclosure; Page 70-71; 133pp; English.
XX
CC The sequence is that of glycerol-3-phosphate dehydrogenase. It was used
CC as part of a method of fermentative production of 1,3-propanediol (1,3-
CC pd), using an organism comprising at least 1 gene encoding a dehydratase,
CC is improved by inserting into the host a gene encoding protein X and
CC culturing the transformant in presence of a carbon source (e.g. mono-,
CC oligo- or poly-saccharide or 1C substrate) convertible to 1,3-pd. 1,3-pd
CC is a starting material for polyesters, polyurethanes and cyclic
CC compounds. 1,3-pd can now be produced by a single recombinant organism
CC from inexpensive carbon sources such as glucose (rather than costly
CC glycerol or dihydroxyacetone), rapidly and without causing pollution
XX
SQ Sequence 339 AA;

Query Match 99.8%; Score 1719; DB 2; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.4e-158;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVFPF 60
DB 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVFPF 60

QY 61 DTLHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
DB 61 DTLHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMRPDARLVWATKGLEAETGRLL 120

QY 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDTQTFADDLQQLLHCGKSPRVY 180
DB 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDTQTFADDLQQLLHCGKSPRVY 180

QY 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRGLGAALGADPATF 240
DB 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRGLGAALGADPATF 240

QY 241 MGMAGLDLVLCTENQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300
DB 241 MGMAGLDLVLCTDNQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300

QY 301 GVEMPIITEEIIYQVLYCGKNAREAAALTLGRARKDERSH 339
DB 301 GVEMPIITEEIIYQVLYCGKNAREAAALTLGRARKDERSH 339

RESULT 4
ID AAY26172 standard; protein; 339 AA.
XX
AC AAY26172;
XX
DT 29-SEP-1999 (first entry)
XX
DE Glycerol-3-phosphate dehydrogenase encoded by gpsA gene.
XX
KW G3PDH; glycerol-3-phosphate dehydrogenase; NADPH-dependent enzyme;
KW EC 1.1.1.94; glycerol; recombinant organism; transformation; gpsA gene;
KW glycerol biosynthetic pathway; expression cassette; 1-3 propanediol;
KW pharmaceutical compound; antifreeze solution; lubricant; polyurethane;
KW cyclic compound; fat and oil industry; polyester fiber.
XX
OS Saccharomyces sp.
XX
PN WO9928480-A1.

XX 10-JUN-1999.
PD
XX
XX 02-DEC-1998; 98WO-US025551.
PF
XX
XX 02-DEC-1997; 97US-00982783.
PR
XX
XX (DUPO) DU PONT DE NEMOURS & CO E I.
PA (GEMV) GENENCOR INT INC.
XX
PI Nair RV, Payne MS, Trimbur DE, Valle F;
XX
XX WPI; 1999-385384/32.
DR
XX
XX Recombinant organisms containing G3PDH and or G3P phosphatase.
PT
XX
XX Claim 12; Page 63-64; 84pp; English.
PS
CC The present sequence is a glycerol-3-phosphate dehydrogenase (G3PDH)
CC enzyme (EC 1.1.1.94) which is NADPH dependent, and catalyses the
CC conversion of dihydroxyacetone phosphate to glycerol-3-phosphate. This is
CC encoded by gps A gene. This is used to produce glycerol from a
CC recombinant organism by transforming a suitable host cell with an
CC expression cassette comprising either one or both of the genes encoding
CC G3PDH and G3P, where the host cell has disruptions in either glycerol
CC kinase or glycerol dehydrogenase endogenous genes to prevent their active
CC expression. The transformed host cell is cultured with a carbon source
CC and glycerol is recovered. Compounds derived from the glycerol
CC biosynthetic pathway like 1,3-propanediol can also be produced. The
CC method provides a rapid, inexpensive and environment-friendly source of
CC glycerol. Glycerol is used in cosmetics, food, pharmaceuticals,
CC lubricants, anti-freeze solutions, fat and oil industry etc. . 1,3 -
CC propanediol is used for the production of polyester fibers and the
CC manufacture of polyurethanes and cyclic compounds
XX
SQ Sequence 339 AA;

Query Match 99.8%; Score 1719; DB 2; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.4e-158;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVFPF 60
DB 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVFPF 60

QY 61 DTLHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
DB 61 DTLHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMRPDARLVWATKGLEAETGRLL 120

QY 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDTQTFADDLQQLLHCGKSPRVY 180
DB 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDTQTFADDLQQLLHCGKSPRVY 180

QY 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRGLGAALGADPATF 240
DB 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRGLGAALGADPATF 240

QY 241 MGMAGLDLVLCTENQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300
DB 241 MGMAGLDLVLCTDNQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300

QY 301 GVEMPIITEEIIYQVLYCGKNAREAAALTLGRARKDERSH 339
DB 301 GVEMPIITEEIIYQVLYCGKNAREAAALTLGRARKDERSH 339

RESULT 5
ID AAU34796 standard; protein; 339 AA.
XX
AC AAU34796;
XX
DT 14-FEB-2002 (first entry)

XX DE E. coli cellular proliferation protein #377.
XX KW Antisense; prokaryotic cellular proliferation protein; antibiotic;
KW antibacterial; drug design.
XX OS Escherichia coli.
XX WO200170955-A2.
XX PD 27-SEP-2001.
XX 21-MAR-2001; 2001WO:US009180.
XX PR 21-MAR-2000; 2000US-0191078P.
PR 23-MAY-2000; 2000US-0206848P.
PR 26-MAY-2000; 2000US-0207727P.
PR 23-OCT-2000; 2000US-0242578P.
PR 27-NOV-2000; 2000US-0253625P.
PR 22-DEC-2000; 2000US-0257931P.
PR 16-FEB-2001; 2001US-0269308P.
XX PA (ELIT-) ELITRA PHARM INC.
XX PI Haselbeck R, Ohlсен KL, Zyskind JW, Wall D, Trawick JD, Carr GJ;
PI Yamamoto RT, Xu HH;
XX DR WPI; 2001-611495/70.
DR N-PSDB; AAS52655.
XX New polynucleotides for the identification and development of
PT antibiotics, comprise sequences of antisense nucleic acids.
XX Example 3; SEQ ID NO 10389; 51lpp; English.
XX The invention relates to antisense inhibitors of genes essential to
CC prokaryotic cellular proliferation, their use in identifying the genes,
CC their use in the discovery of novel antibiotics, the essential genes
CC themselves and the encoded proteins. The prokaryotes used are Escherichia
CC coli, Staphylococcus aureus, Salmonella typhi, Klebsiella pneumoniae,
CC Pseudomonas aeruginosa and Enterococcus faecalis. The invention is also
CC useful for the identification of potential new targets for antibiotic
CC development. The antisense nucleic acids can also be used to identify
CC proteins used in proliferation, to express these proteins, and to obtain
CC antibodies capable of binding to the expressed proteins. The proteins can
CC be used to screen compounds in rational drug discovery programmes. The
CC antisense nucleic acid sequence is also useful to screen for homologous
CC nucleic acids which are required for cell proliferation in a wide variety
CC of organisms. The present sequence represents an essential prokaryotic
CC cellular proliferation protein. Note: The sequence data for this patent
CC did not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX SQ Sequence 339 AA;
Query Match 99.8%; Score 1719; DB 4; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.4e-158;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPPF 60
Db 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPPF 60
QY 61 DTLHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMRPDLARLVWATKGLAEATGRLL 120
Db 61 DTLHLESDLATALAASRNILVVVPSHVFGVLRQIKPLMRPDLARLVWATKGLAEATGRLL 120
QY 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLHCGKSFVY 180
Db 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLHCGKSFVY 180
QY 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLAEMSRLGAALGADPATF 240

Db 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLAEMSRLGAALGADPATF 240
QY 241 MGMAGLGDLVLTCTENQSRNRRFGMMLGQGMVDVQSAQEKIGVVVEGYRNTKEVRELAHRF 300
Db 241 MGMAGLGDLVLTCTDNQSRNRRFGMMLGQGMVDVQSAQEKIGVVVEGYRNTKEVRELAHRF 300
QY 301 GVEMPTITEEIVQVLYCGKNAREAAALTLLGRARKDERSH 339
Db 301 GVEMPTITEEIVQVLYCGKNAREAAALTLLGRARKDERSH 339
RESULT 6
ABU28819
ID ABU28819 standard; protein; 339 AA.
XX AC ABU28819;
XX DT 19-JUN-2003 (first entry)
XX DE Protein encoded by Prokaryotic essential gene #14346.
XX KW Antisense; prokaryotic essential gene; cell proliferation; drug design.
XX OS Escherichia coli.
XX PN WO200277183-A2.
XX PD 03-OCT-2002.
XX 21-MAR-2002; 2002WO-US009107.
PF 21-MAR-2001; 2001US-00815242.
XX PR 06-SEP-2001; 2001US-00948993.
PR 25-OCT-2001; 2001US-0342923P.
PR 08-FEB-2002; 2002US-00072851.
PR 06-MAR-2002; 2002US-0362699P.
XX PA (ELIT-) ELITRA PHARM INC.
XX PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlсен KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
XX WPI; 2003-029926/02.
DR N-PSDB; ACA32689.
XX New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.
PS Claim 25; SEQ ID NO 56743; 1766pp; English.
XX The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the

CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is encoded by one of
CC the target prokaryotic essential genes. Note: The sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 339 AA;

Query Match 99.8%; Score 1719; DB 6; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.4e-158;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60
Db 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60

QY 61 DTLHLESDLATALAASRNILVVVPSHVFEVLRLQIKPLMRPDARLVWATKGLEAETGRLL 120
Db 61 DTLHLESDLATALAASRNILVVVPSHVFEVLRLQIKPLMRPDARLVWATKGLEAETGRLL 120

QY 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLHCGKSFVRV 180
Db 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLHCGKSFVRV 180

QY 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATF 240
Db 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATF 240

QY 241 MGMAGLDLVLTCTENQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300
Db 241 MGMAGLDLVLTCTENQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300

QY 301 GVEMPTIEEIIQVLYCGKNAREAAALTLGRARKDERSH 339
Db 301 GVEMPTIEEIIQVLYCGKNAREAAALTLGRARKDERSH 339

RESULT 7
ADS45174
ID ADS45174 standard; protein; 339 AA.
XX
AC ADS45174;
XX
DT 02-DEC-2004 (first entry)
XX
DE Bacterial polypeptide #23604.
XX
KW Recombinant DNA construct; transformed plant; improved plant property;
KW cold tolerance; heat tolerance; drought tolerance; herbicide; osmosis;
KW pathogen tolerance; pest tolerance; plant disease resistance;
KW cell cycle pathway modification; plant growth regulator;
KW homologous recombination; seed oil yield; protein yield; carbohydrate;
KW nitrogen; phosphorus; photosynthesis; lignin; galactomannan;
KW bacterial polypeptide.
XX
OS Bacteria.
XX
PN US2003233675-A1.
XX
PD 18-DEC-2003.
XX
PF 20-FEB-2003; 2003US-00369493.
XX
PR 21-FEB-2002; 2002US-0360039P.
XX
PA (CAOY/) CAO Y.
PA (HINK/) HINKLE G J.
PA (SLAT/) SLATER S C.

PA (CHEN/) CHEN X.
PA (GOLD/) GOLDMAN B S.
XX
PI Cao Y, Hinkle GJ, Slater SC, Chen X, Goldman BS;
XX WPI; 2004-061375/06.
DR
XX
PT New recombinant DNA construct comprising a promoter positioned to provide
PT for expression of a polynucleotide encoding a polypeptide from a
PT microbial source, useful for producing plants with improved properties.
XX
PS Claim 1; SEQ ID NO 23604; 122pp; English.
XX
CC The invention relates to a recombinant DNA construct comprising a
CC promoter functional in a plant cell, where the promoter is positioned to
CC provide for expression of a polynucleotide encoding a polypeptide from a
CC microbial source. The invention also relates to a transformed plant
CC comprising the recombinant DNA construct and a method of producing a
CC transformed plant having an improved property. The plant is a crop plant
CC such as maize or soybean. The method of producing a transformed plant
CC having an improved property comprises transforming a plant with the
CC recombinant DNA construct and growing the transformed plant, where the
CC polynucleotide or polypeptide is useful for improving plant properties.
CC The recombinant DNA construct is useful for producing plants with
CC improved plant properties, e.g. improved cold, heat or drought tolerance,
CC tolerance to herbicides, extreme osmotic conditions, pathogens or pests,
CC increased resistance to plant disease, better growth rate by modification
CC of the cell cycle pathway with plant growth regulators, increased rate of
CC homologous recombination, modified seed oil or protein yield and/or
CC content, improved yield by modification of carbohydrate, nitrogen or
CC phosphorus use and/or uptake, by modification of photosynthesis or by
CC providing improved plant growth and development under at least one stress
CC condition, improved lignin production or improved galactomannan
CC production. This sequence represents a bacterial polypeptide used in the
CC scope of the invention. Note: The sequence data for this patent did not
CC form part of the printed specification but was obtained in electronic
CC format from USPTO at seqdata.uspto.gov/sequence.html.
XX
SQ Sequence 339 AA;

Query Match 99.8%; Score 1719; DB 8; Length 339;
Best Local Similarity 99.7%; Pred. No. 1.4e-158;
Matches 338; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60
Db 1 MNQRNASMTVIGAGSYGTALAITLARNGHEVVLWGHDPHEIATLERDRCNAAFLPDVPFP 60

QY 61 DTLHLESDLATALAASRNILVVVPSHVFEVLRLQIKPLMRPDARLVWATKGLEAETGRLL 120
Db 61 DTLHLESDLATALAASRNILVVVPSHVFEVLRLQIKPLMRPDARLVWATKGLEAETGRLL 120

QY 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLHCGKSFVRV 180
Db 121 QDVAREALGDQIPLAVISGPTFAKELAAGLPTAISLASTDQTFADDLQQLHCGKSFVRV 180

QY 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATF 240
Db 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGI GFGANARTALITRGLAEMSRLGAALGADPATF 240

QY 241 MGMAGLDLVLTCTENQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300
Db 241 MGMAGLDLVLTCTENQSRNRRFGMMLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRF 300

QY 301 GVEMPTIEEIIQVLYCGKNAREAAALTLGRARKDERSH 339
Db 301 GVEMPTIEEIIQVLYCGKNAREAAALTLGRARKDERSH 339

RESULT 8
ABU47465
ID ABU47465 standard; protein; 339 AA.
XX

PT New nucleic acid encoding a Klebsiella pneumoniae polypeptide, useful for
PT preparing a vaccine composition against Klebsiella pneumoniae.

XX Disclosure; SEQ ID NO 10740; 932pp; English.

XX The invention describes a new isolated nucleic acid encoding a Klebsiella
CC pneumoniae polypeptide. Also described are: a recombinant expression
CC vector comprising the nucleic acid, operably linked to a transcription
CC regulatory element; and a cell comprising the recombinant expression
CC vector. The nucleic acid is useful for preparing a vaccine composition
CC against Klebsiella pneumoniae. This is the amino acid sequence of a
CC Klebsiella pneumoniae polypeptide of the invention

XX Sequence 345 AA;

SQ Query Match 92.2%; Score 1587; DB 7; Length 345;
Best Local Similarity 92.3%; Pred. No. 1.1e-145;
Matches 313; Conservative 11; Mismatches 15; Indels 0; Gaps 0;

Qy 1 MNQRNASMTVIGAGSYGTALAITLARNHGHEVVLWGHDPKHATLERDRCNAFLPDVPPF 60
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
7 MNALNAAMTVIGAGSYGTALAITLARNGHVVLWGHDPKHATLQHDRCNAFLPDVPPF 66
Qy 61 DTLHLESDLTALAASRNILVVVPSHVFGVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
67 DTLHLESDLTALAASRDILVVVPSHVFGVLRQIKPLMRSDARLVWATKGLEAETGRLL 126
Qy 121 QDVAREALGDOIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLLHCGKSRVY 180
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
127 QDVAREALGDDIPLAVISGPTFAKELAAAGLPTAISLAATDPQFAEDLQRLLLHCGKSRVY 186
Qy 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLAEMSRLGAALGADPATF 240
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
187 INPDFIGVQLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLVEMSRGLAALGADPETF 246
Qy 241 MGMAGLGDVLVLTCTENQSRNRRFGMMLGQGMVDVQSAQDKIGQVVEGYRNTKEVRELAHRF 300
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
247 MGMAGLGDVLVLTCTDNQSRNRRFGMMLGQGMVDVQSAQDKIGQVVEGYRNTKEVRLAQRL 306
Qy 301 GVEMPIITEIYQVLYCGKNAREAAALTLGRARKDERSH 339
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
307 GVEMPIITEIYQVLYCGKIAREAAALTLGRARKDERSN 345

RESULT 13
ABU50149
ID ABU50149 standard; protein; 339 AA.
XX AC ABU50149;
XX 19-JUN-2003 (first entry)
XX Protein encoded by Prokaryotic essential gene #35676.
XX Antisense; prokaryotic essential gene; cell proliferation; drug design.
XX Yersinia pestis.
XX WO200277183-A2.
XX 03-OCT-2002.
XX 21-MAR-2002; 2002WO-US009107.
XX 21-MAR-2001; 2001US-00815242.
PR 06-SEP-2001; 2001US-00948993.
PR 25-OCT-2001; 2001US-0342923P.
PR 08-FEB-2002; 2002US-00072851.
PR 06-MAR-2002; 2002US-0362699P.
XX (ELIT-) ELITRA PHARM INC.
XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;

PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
XX WPI; 2003-029926/02.
DR N-PSDB; ACA54019.

XX New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.

PS Claim 25; SEQ ID NO 78073; 1766pp; English.

XX The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than S. aureus, S. typhimurium,
CC K. pneumoniae or P. aeruginosa. The present sequence is encoded by one of
CC the target prokaryotic essential genes. Note: The sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX

SQ Sequence 339 AA;

Query Match 84.5%; Score 1455; DB 6; Length 339;
Best Local Similarity 84.2%; Pred. No. 7.5e-133;
Matches 283; Conservative 24; Mismatches 29; Indels 0; Gaps 0;

Qy 1 MNQRNASMTVIGAGSYGTALAITLARNHGHEVVLWGHDPKHATLERDRCNAFLPDVPPF 60
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
1 MNTNPASMAVIGAGSYGTALAITLARNGHQVVLWGHDPKHATLQHDRCNRAFLPDAAFP 60
Qy 61 DTLHLESDLTALAASRNILVVVPSHVFGVLRQIKPLMRPDARLVWATKGLEAETGRLL 120
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
61 DTLRLETDLACALAASRDVLVVVPSHVFGAVLHQLKPHLRKDARIVWATKGLEAETGRLL 120
Qy 121 QDVAREALGDOIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLLHCGKSRVY 180
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
121 QDVAREVLGEAIPAVISGPTFAKELAAAGLPTAIALASTDVFQSEDLQQLLHCGKSRVY 180
Qy 181 SNPDFIGVQLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLAEMSRLGAALGADPATF 240
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
181 SNPDFIGVQLGGAVKNVIAIGAGMSDGIGFGANARTALITRGLAEMTRLTALGADPSTF 240
Qy 241 MGMAGLGDVLVLTCTENQSRNRRFGMMLGQGMVDVQSAQDKIGQVVEGYRNTKEVRELAHRF 300
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
241 MGMAGLGDVLVLTCTDNQSRNRRFGIMLGQGLGVKEAQDNIGQVVEGYRNTKEVLAQRH 300
Qy 301 GVEMPIITEIYQVLYCGKNAREAAALTLGRARKDER 336
Db |||:|||||||||||||||||||||||||||||||||||||||||||||||||||||||||
301 GVEMPIITEIYQVLYCHKNAREAAALTLGRTKKDEK 336

CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is encoded by one of
CC the target prokaryotic essential genes. Note: The sequence data for this
CC patent did not form part of the printed specification, but was obtained
CC in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published/pct/sequences

Query Match	81.9%;	Score 1411;	DB 6;	Length 337;
Best Local Similarity	82.0%;	Pred. No. 1.5e-128;		
Matches 274;	Conservative	27;	Mismatches 33;	Indels 0; Gaps 0;
QY	5	NASMTVIGAGSYGTALAITLARNGHEWVLWGHDPHEIATLERDRCNAAFLPDVPFPDTLH	64	
Db	2	NASMTVIGAGSYGTALAITLARNGHDVVLWGHDPKHVAALDEQARCNQAFLPDVSFPDSLY	61	
QY	65	LESDLATALAASRNILVVVPSHVFEVLROIKPLMRPDARLWVATKGLEAETGRLLQDVA	124	
Db	62	MEASLQKAIEASRNILVVPISHVFEVLQOIKPFLRQDARVVWATKGLEAHTGRLLQDVA	121	
QY	125	REALGDQIPLAVISGPTFAKELAAAGLPTAISLASTDQTFADDLQQLHCGKSFVYSNPD	184	
Db	122	REVLGNEIPLAVLSGPTFAKELAAAGLPTAIAVASTDNLFLQLQQLFHCCKSFVYKNPD	181	
QY	185	FIGVQLGGAVKNVIAIGAGMSDCIGFGANARTALITRGLAEMSRLLGAALGADPATFMGVA	244	
Db	182	FIGVQLGGAVKNVIAIGAGMSDGMGFGANARTALITRGLAEMSRLLGKALGADAATFMGVA	241	
QY	245	GLGDLVLTCTENQSRNRRFGMWLGQGMVDVQSAQEKIGQVVEGYRNTKEVRELAHRFVEM	304	
Db	242	GLGDLVLTCTDNQSRNRRFGMWLGQGFVDTAQEKIGQVVEGYRNTKEVRALAEQVGVEM	301	
QY	305	PITEEIQVLYCGKNAREAAITLLGRARKDERS	338	
Db	302	PITEEIQVILYQHKVDVKEAALALLGRATKDEIDS	335	

Search completed: April 27, 2005, 10:59:44
Job time : 167 secs

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OM nucleic - nucleic search, using sw model

Run on: April 27, 2005, 09:52:56 ; Search time 656 Seconds
(without alignments)
9204.476 Million cell updates/sec

Title: US-10-088-079-1
Perfect score: 1020
Sequence: 1 atgaaccaacgtaagtcttc.....acgagcgagcagccactaa 1020

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0
Maximum DB seq length: 20000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N Geneseq_16Dec04:*
1: geneseqn1980s:*
2: geneseqn1990s:*
3: geneseqn2000s:*
4: geneseqn2001as:*
5: geneseqn2001bs:*
6: geneseqn2002as:*
7: geneseqn2002bs:*
8: geneseqn2003as:*
9: geneseqn2003bs:*
10: geneseqn2003cs:*
11: geneseqn2003ds:*
12: geneseqn2004as:*
13: geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	1020	100.0	1020	5	Aaf57428 E. coli g
2	1018.4	99.8	1020	4	Aas52655 E. coli D
3	1018.4	99.8	1020	8	Aca32689 Prokaryot
4	1018.4	99.8	1020	13	Adt48853 Bacterial
5	787	77.2	1020	8	Aca51335 Prokaryot
6	765.8	75.1	1017	8	Aca31966 Prokaryot
7	756.8	74.2	1017	8	Aca35838 Prokaryot
8	756.6	74.2	1038	11	Ach97774 Klebsiell
9	749.4	73.5	1023	8	Aca49224 Prokaryot
10	606	59.4	1020	8	Aca54019 Prokaryot
11	565.6	55.5	781	6	Abq21987 Oligonucl
12	565.6	55.5	781	6	Abq21986 Oligonucl
13	556.4	54.5	1023	10	Acf70491 Photorhab
14	556.4	54.5	69727	10	Acf65374 Photorhab
15	556.4	54.5	110000	10	Acf67367_35
16	539.6	52.9	1011	8	Aca44463 Prokaryot
17	539.6	52.9	1023	10	Adf03046 Bacterial
18	539.4	52.9	990	13	Ads45613 Bacterial
19	521.2	51.1	990	13	Adt46572 Bacterial
20	513.8	50.4	781	6	Abq21989 Oligonucl

C	21	513.8	50.4	781	6	ABQ21988	Abq21988 Oligonucl
	22	476.2	46.7	1035	8	ACA53490	Aca53490 Prokaryot
	23	414.4	40.6	1014	8	ACA43241	Aca43241 Prokaryot
	24	399.4	39.2	135356	13	ADT05646	Adt05646 Haemophil
	25	396.2	38.8	1008	4	AAS53331	Aas53331 Haemophil
	26	396.2	38.8	1008	8	ACA34182	Aca34182 Prokaryot
	27	396.2	38.8	110000	2	AAT42063_06	Continuation (7 of
C	28	388.4	38.1	4711	13	ADT05428	Adt05428 Haemophil
	29	325.6	31.9	503	6	ABQ50341	Abq50341 Oligonucl
C	30	325.6	31.9	503	6	ABQ50340	Abq50340 Oligonucl
C	31	313.8	30.8	503	6	ABQ50339	Abq50339 Oligonucl
	32	313.8	30.8	503	6	ABQ50338	Abq50338 Oligonucl
	33	238.6	23.4	1002	13	ADT41591	Adt41591 Bacterial
	34	238.6	23.4	1002	13	ADS63997	Ads63997 Bacterial
	35	238.6	23.4	1020	13	ADS63622	Ads63622 Bacterial
	36	221	21.7	1041	13	ADT42864	Adt42864 Bacterial
	37	219.4	21.5	987	8	ACA37060	Aca37060 Prokaryot
	38	211.4	20.7	1002	13	ADS57169	Ads57169 Bacterial
	39	199.8	19.6	987	8	ABX09916	Abx09916 N. mening
	40	199.8	19.6	49767	3	AAA81458	Aaa81458 N. mening
C	41	199.8	19.6	110000	3	AAA81489_7	Continuation (8 of
C	42	199.8	19.6	110000	4	AAI99682_33	Continuation (34 o
C	43	199.8	19.6	110000	4	AAI99683_33	Continuation (34 o
C	44	199.8	19.6	172325	3	AAF21613	Aaf21613 Neisseria
	45	199.6	19.6	1002	8	ACA38486	Aca38486 Prokaryot

ALIGNMENTS

RESULT 1
AAF57428
ID AAF57428 standard; DNA; 1020 BP.

XX AAF57428;

DT 11-JUN-2001 (first entry)

XX E. coli gpsA2FR encoding DNA.

KW Glycerol-3-phosphate dehydrogenase; G3PD; feedback inhibition; oil seed;
KW genetic transformation; fatty acid; glycerolipid; osmotic stress; gpsA;
KW gpsA2FR; allele; ds.

OS Escherichia coli.

XX Key Location/Qualifiers
FT CDS 1..1020

FT mutation /tag= a
FT /product= "gpsA2FR"

FT /tag= b
FT /note= "there is a point mutation at this position as compared to the wild-type gpsA gene, which makes the gene feed-defective; wild-type GAC codon is changed to GAA codon"

PN WO200121820-A1.

PD 29-MAR-2001.

PF 21-SEP-2000; 2000WO-CA001096.

XX 22-SEP-1999; 99US-0155133P.

XX (CANA) NAT RES COUNCIL CANADA.

XX Zou J, Wei Y, Periappuram C, Selvaraj G, Datla R;

DR WPI; 2001-257996/26.

DR P-PSDB; AAB62189.

PT Manipulating glycerol-3-phosphate metabolism of plant for enhancing

PT stress tolerance, altering fatty acid content in glycerolipids, by
PT expressing in plant feedback defective glycerol-3-phosphate dehydrogenase
PT gene.
XX

XX Claim 5; Fig 1; 39pp; English.

PS The invention provides a method for genetically transforming a plant so
XX that it expresses a heterologous glycerol-3-phosphate dehydrogenase
CC (G3PD) that is less sensitive to feedback inhibition than wild-type G3PD.
CC The method involves providing a vector comprising a DNA sequence encoding
CC G3PD that is less sensitive to feedback inhibition than wild-type G3PD
CC and transforming the plant with the vector. The method is useful for
CC expressing a heterologous G3PD less sensitive to feedback inhibition than
CC wild-type G3PD in an oil seed bearing plant, such as Arabidopsis thaliana
CC or Brassica. The vectors are useful for producing a genetically altered
CC plant having altered fatty acid content in its glycerolipids, especially
CC elevated levels of C16 fatty acids and increased osmotic stress tolerance
CC relative to the wild type. The present sequence represents the DNA
CC encoding the E. coli gpsA2FR protein. The gene gpsA2FR is an allele of
CC the E. coli gpsA gene, and encodes an altered version of the GPDH protein
CC defective in feedback inhibition. This gpsA2FR gene can be used in the
CC vectors and method of the invention
XX

SQ Sequence 1020 BP; 214 A; 274 C; 304 G; 228 T; 0 U; 0 Other;

Query Match 100.0%; Score 1020; DB 5; Length 1020;
Best Local Similarity 100.0%; Pred. No. 2.1e-308;
Matches 1020; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCCGGCTCGTACGGCACCGTCTTT 60
Db |
1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCCGGCTCGTACGGCACCGTCTTT 60
QY 61 GCCATCACCTTGGCAAGAAATGGCCACGAGTGTCTCTGGGGCCATGACCCTGAACAT 120
Db |
61 GCCATCACCTTGGCAAGAAATGGCCACGAGTGTCTCTGGGGCCATGACCCTGAACAT 120
QY 121 ATCGCAACGCTTGAACGCGACCGCTGTAAACGCGCGTCTTCTCCCGATGTGCTTTTCCC 180
Db |
121 ATCGCAACGCTTGAACGCGACCGCTGTAAACGCGCGTCTTCTCCCGATGTGCTTTTCCC 180
QY 181 GATACGCTCCATCTTGAAAGCGATCTCGCCACTCGCTGGCAGCCAGCCGTAATATCTC 240
Db |
181 GATACGCTCCATCTTGAAAGCGATCTCGCCACTCGCTGGCAGCCAGCCGTAATATCTC 240
QY 241 GTCGTCGTACCCAGCCATGTCTTTGGTGAAGTCTGCGCCAGATTAAACCACTGATCGGT 300
Db |
241 GTCGTCGTACCCAGCCATGTCTTTGGTGAAGTCTGCGCCAGATTAAACCACTGATCGGT 300
QY 301 CCTGATCGCGTCTGGTGTGGCGACCAAGGGCTGGAAGCGGAAACCGGACGTCTGTTA 360
Db |
301 CCTGATCGCGTCTGGTGTGGCGACCAAGGGCTGGAAGCGGAAACCGGACGTCTGTTA 360
QY 361 CAGGACGTGGCGGTGAGGCCCTTAGCGGATCAAAATCCGCTGGCGGTTATCTCTGGCCCA 420
Db |
361 CAGGACGTGGCGGTGAGGCCCTTAGCGGATCAAAATCCGCTGGCGGTTATCTCTGGCCCA 420
QY 421 ACGTTTGCGAAAGAACTGGCGGAGGTTTACCGACAGCTATTTTCGCTGGCTCGACCGAT 480
Db |
421 ACGTTTGCGAAAGAACTGGCGGAGGTTTACCGACAGCTATTTTCGCTGGCTCGACCGAT 480
QY 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACGAGTTTACCGACAGCTATTTTCGCTGGCTCGACCGAT 540
Db |
481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACGAGTTTACCGACAGCTATTTTCGCTGGCTCGACCGAT 540
QY 541 AGCAATCCGGATTTCATTGGCGTGCAGCTTGGCGGCGGTTGAAACCGTTATTGCCATT 600
Db |
541 AGCAATCCGGATTTCATTGGCGTGCAGCTTGGCGGCGGTTGAAACCGTTATTGCCATT 600
QY 601 GGTGCGGGGATGTCGACGGTATCGGTTTGGTGCGAATGCGCGTACGGCGCTGATCACC 660
Db |
601 GGTGCGGGGATGTCGACGGTATCGGTTTGGTGCGAATGCGCGTACGGCGCTGATCACC 660

QY 661 CGTGGGCTGGCTGAAATGTGCGGCTCTTGTCGGCGCTGGGTGCCGACCCCTGCCACCTTT 720
Db |
661 CGTGGGCTGGCTGAAATGTGCGGCTCTTGTCGGCGCTGGGTGCCGACCCCTGCCACCTTT 720
QY 721 ATGGGCATGGCGGGCTTGGCGATCTGGTGCTTACCTGTACCGAAAAACAGTCGCGTAAC 780
Db |
721 ATGGGCATGGCGGGCTTGGCGATCTGGTGCTTACCTGTACCGAAAAACAGTCGCGTAAC 780
QY 781 CGCCGTTTTGGCATGCTCGGTCAAGGCATGGATGTACAAAGCGCGCAGGAGAAGATT 840
Db |
781 CGCCGTTTTGGCATGCTCGGTCAAGGCATGGATGTACAAAGCGCGCAGGAGAAGATT 840
QY 841 GGTCAGGTGCTGGAAGGCTACCGCAATACGAAAGAAAGTCCGCGAACTGCGGCATCGCTTC 900
Db |
841 GGTCAGGTGCTGGAAGGCTACCGCAATACGAAAGAAAGTCCGCGAACTGCGGCATCGCTTC 900
QY 901 GGCGTTGAATGCCAATAACCGAGGAATTTATCAAGTATTATATTCGGAAAAACGCG 960
Db |
901 GGCGTTGAATGCCAATAACCGAGGAATTTATCAAGTATTATATTCGGAAAAACGCG 960
QY 961 CGCGAGGCGACATTGACTTTACTAGTTCGTGCACGCAAGGACGAGCGCAGCCACTAA 1020
Db |
961 CGCGAGGCGACATTGACTTTACTAGTTCGTGCACGCAAGGACGAGCGCAGCCACTAA 1020

RESULT 2

AAS52655

ID AAS52655 standard; DNA; 1020 BP.

XX AAS52655;

DT 13-FEB-2002 (first entry)

XX E. coli DNA for cellular proliferation protein #377.

DE Antisense; ds; prokaryotic cellular proliferation gene; antibiotic;

XX antibacterial; drug design.

OS Escherichia coli.

XX WO200170955-A2.

PN 27-SEP-2001.

PD 21-MAR-2001; 2001WO-US009180.

XX 21-MAR-2000; 2000US-0191078P.

PR 23-MAY-2000; 2000US-0206848P.

PR 26-MAY-2000; 2000US-0207727P.

PR 23-OCT-2000; 2000US-0242578P.

PR 27-NOV-2000; 2000US-0253625P.

PR 22-DEC-2000; 2000US-0257931P.

PR 16-FEB-2001; 2001US-0269308P.

XX (ELIT-) ELITRA PHARM INC.

XX Haselbeck R, Ohlsen KL, Zyskind JW, Wall D, Trawick JD, Carr GJ;

PI Yamamoto RT, Xu HH;

XX WPI; 2001-611495/70.

DR P-PSDB; AAU34796.

XX New polynucleotides for the identification and development of

PT antibiotics, comprise sequences of antisense nucleic acids.

XX Claim 27; SEQ ID NO 6292; 511pp; English.

CC The invention relates to antisense inhibitors of genes essential to

CC prokaryotic cellular proliferation, their use in identifying the genes,

CC their use in the discovery of novel antibiotics, the essential genes

CC themselves and the encoded proteins. The prokaryotes used are Escherichia

CC coli, Staphylococcus aureus, Salmonella typhi, Klebsiella pneumoniae,

CC Pseudomonas aeruginosa and Enterococcus faecalis. The invention is also

CC useful for the identification of potential new targets for antibiotic
CC development. The antisense nucleic acids can also be used to identify
CC proteins used in proliferation, to express these proteins, and to obtain
CC antibodies capable of binding to the expressed proteins. The proteins can
CC be used to screen compounds in rational drug discovery programmes. The
CC antisense nucleic acid sequence is also useful to screen for homologous
CC nucleic acids which are required for cell proliferation in a wide variety
CC of organisms. The present sequence encodes an essential prokaryotic
CC cellular proliferation protein. Note: The sequence data for this patent
CC did not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX

SQ Sequence 1020 BP; 213 A; 275 C; 304 G; 228 T; 0 U; 0 Other;

Query Match 99.8%; Score 1018.4; DB 4; Length 1020;
Best Local Similarity 99.9%; Pred. No. 6.8e-308;
Matches 1019; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTCCGGCTCGTACGGCACCCTCTT 60
DB 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTCCGGCTCGTACGGCACCCTCTT 60
QY 61 GCCATCACCTGGCAAGAAATGGCCACGAGGTGTCTCTGGGGCCATGACCTGAACAT 120
DB 61 GCCATCACCTGGCAAGAAATGGCCACGAGGTGTCTCTGGGGCCATGACCTGAACAT 120
QY 121 ATCGCAACGCTTGAAACGCGACCGCTGTAAACGCCCGTCTTCTCCCGATGTGCTTCCC 180
DB 121 ATCGCAACGCTTGAAACGCGACCGCTGTAAACGCCCGTCTTCTCCCGATGTGCTTCCC 180
QY 181 GATACGCTCCATCTTGAAAGCGATCTCGCCACTCGGCTGGCAGCCAGCCGTAATATCTC 240
DB 181 GATACGCTCCATCTTGAAAGCGATCTCGCCACTCGGCTGGCAGCCAGCCGTAATATCTC 240
QY 241 GTCGTGTAACCCAGCCATGCTTTGGTGAAGTGTGCGCCAGATTAAACCACTGATCGGT 300
DB 241 GTCGTGTAACCCAGCCATGCTTTGGTGAAGTGTGCGCCAGATTAAACCACTGATCGGT 300
QY 301 CCTGATGCGCGTCTGCTGGGCGACCAAGGGCTGGAAGCGGAAACCGGACGTCTGTTA 360
DB 301 CCTGATGCGCGTCTGCTGGGCGACCAAGGGCTGGAAGCGGAAACCGGACGTCTGTTA 360
QY 361 CAGGACGTGGCGGTGAGGCCCTTAGCGGATCAAAATCCGCTGGCGGTTATCTTGGCCCA 420
DB 361 CAGGACGTGGCGGTGAGGCCCTTAGCGGATCAAAATCCGCTGGCGGTTATCTTGGCCCA 420
QY 421 ACGTTTGCAGAAAGAACTGGCGGCGAGGTTTACCGACAGCTATTTTCGTGGCCTCGACCGAT 480
DB 421 ACGTTTGCAGAAAGAACTGGCGGCGAGGTTTACCGACAGCTATTTTCGTGGCCTCGACCGAT 480
QY 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGCGGCAAAAGTTTCCGCGTTAC 540
DB 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGCGGCAAAAGTTTCCGCGTTAC 540
QY 541 AGCAATCCGGATTTCATTTGGCGTGCAGCTTGGCGGCGGCTGAAACACGTTATTGCCATT 600
DB 541 AGCAATCCGGATTTCATTTGGCGTGCAGCTTGGCGGCGGCTGAAACACGTTATTGCCATT 600
QY 601 GGTGCGGGGATGTCGACCGGTATCGGTTTGGTGGGAATGCGGTACGGCGGTGATCACC 660
DB 601 GGTGCGGGGATGTCGACCGGTATCGGTTTGGTGGGAATGCGGTACGGCGGTGATCACC 660
QY 661 CGTGGGCTGGTGAATGTCGCGTCTTGGTGGCGGCGTGGGTGCCGACCTTTT 720
DB 661 CGTGGGCTGGTGAATGTCGCGTCTTGGTGGCGGCGTGGGTGCCGACCTTTT 720
QY 721 ATGGGCATGGCGGGCTTGGCGATCTGGTGTCTTACCTGTACCGAAACACGTCGCGTAAC 780
DB 721 ATGGGCATGGCGGGCTTGGCGATCTGGTGTCTTACCTGTACCGAAACACGTCGCGTAAC 780
QY 781 CGCCGTTTGGCATGATGCTCGGTGAGGGCATGGATGTACAAAGCGCGCAGGAGGATT 840
DB 781 CGCCGTTTGGCATGATGCTCGGTGAGGGCATGGATGTACAAAGCGCGCAGGAGGATT 840

Db 781 CGCCGTTTGGCATGATGCTCGGTGAGGGCATGGATGTACAAAGCGCGCAGGAGGATT 840
QY 841 GGTGAGGTGGTGAAGGCTACCGCAATACGAAAGAAAGTCCGGAACCTGGCGATCGCTTC 900
Db 841 GGTGAGGTGGTGAAGGCTACCGCAATACGAAAGAAAGTCCGGAACCTGGCGATCGCTTC 900
QY 901 GGCCTTGAATGCAATGCAATGCAATGCAATGCAATGCAATGCAATGCAATGCAATGCAAT 960
Db 901 GGCCTTGAATGCAATGCAATGCAATGCAATGCAATGCAATGCAATGCAATGCAATGCAAT 960
QY 961 CGCGAGGCGAGCATTTGACTTTTACTAGGTCGTGCGCAAGGACGCGCAGCAGCACTAA 1020
Db 961 CGCGAGGCGAGCATTTGACTTTTACTAGGTCGTGCGCAAGGACGCGCAGCAGCACTAA 1020

RESULT 3

ACA32689

ID ACA32689 standard; DNA; 1020 BP.

XX AC ACA32689;

XX DT 19-JUN-2003 (first entry)

XX DE Prokaryotic essential gene #14346.

XX KW Antisense; ds; prokaryotic essential gene; cell proliferation;
XX drug design; gene.

XX OS Escherichia coli.

XX PN WO200277183-A2.

XX PD 03-OCT-2002.

XX PF 21-MAR-2002; 2002WO-US009107.

XX PR 21-MAR-2001; 2001US-00815242.

XX PR 06-SEP-2001; 2001US-00948993.

XX PR 25-OCT-2001; 2001US-0342923P.

XX PR 08-FEB-2002; 2002US-00072851.

XX PR 06-MAR-2002; 2002US-0362699P.

XX PA (ELIT-) ELITRA PHARM INC.

XX PI Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;

XX PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;

XX DR WPI; 2003-029926/02.

XX DR P-PSDB; ABU28819.

XX PT New antisense nucleic acids, useful for identifying proteins or screening
XX for homologous nucleic acids required for cellular proliferation to
XX isolate candidate molecules for rational drug discovery programs.

XX PS Claim 14; SEQ ID NO 20559; 1766pp; English.

XX CC The invention relates to an isolated nucleic acid comprising any one of
XX the 6213 antisense sequences given in the specification where expression
XX of the nucleic acid inhibits proliferation of a cell. Also included are:
XX (1) a vector comprising a promoter operably linked to the nucleic acid
XX encoding a polypeptide whose expression is inhibited by the antisense
XX nucleic acid; (2) a host cell containing the vector; (3) an isolated
XX polypeptide or its fragment whose expression is inhibited by the
XX antisense nucleic acid; (4) an antibody capable of specifically binding
XX the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
XX proliferation or the activity of a gene in an operon required for
XX proliferation; (7) identifying a compound that influences the activity of
XX the gene product or that has an activity against a biological pathway
XX required for proliferation, or that inhibits cellular proliferation; (8)
XX identifying a gene required for cellular proliferation or the biological
XX pathway in which a proliferation-required gene or its gene product lies
XX or a gene on which the test compound that inhibits proliferation of an
XX organism acts; (9) manufacturing an antibiotic; (10) profiling a

CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
CC prokaryotic essential genes. Note: The sequence data for this patent did
CC not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX

SQ Sequence 1020 BP; 213 A; 275 C; 304 G; 228 T; 0 U; 0 Other;

Query Match 99.8%; Score 1018.4; DB 8; Length 1020;
Best Local Similarity 99.9%; Pred. No. 6.8e-308;
Matches 1019; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY	1	ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTCCGGCTCGTACGACACCGCTCTT	60
Db	1	ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTCCGGCTCGTACGACACCGCTCTT	60
QY	61	GCCATCACCTGGCAAGAAATGGCCACGAGTTGTCTCTCTGGGGCCATGACCCTGAACAT	120
Db	61	GCCATCACCTGGCAAGAAATGGCCACGAGTTGTCTCTCTGGGGCCATGACCCTGAACAT	120
QY	121	ATCGCAACGCTTGAACCGGACCGCTGTAAACGCCGCGTTTCTCCCGATGTGCTTTCCC	180
Db	121	ATCGCAACGCTTGAACCGGACCGCTGTAAACGCCGCGTTTCTCCCGATGTGCTTTCCC	180
QY	181	GATACGCTCCATCTTTGAAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAATATCTC	240
Db	181	GATACGCTCCATCTTTGAAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAATATCTC	240
QY	241	GTCGTCGTACCCAGCCATGTCTTTGGTGAAGTGTGCGCCAGATTAAACCACTGATGCGT	300
Db	241	GTCGTCGTACCCAGCCATGTCTTTGGTGAAGTGTGCGCCAGATTAAACCACTGATGCGT	300
QY	301	CCTGATCGCGCTCTGGTGTGGCGACCAAGSGCTGGAAGCGAAACCGGACGTCTGTTA	360
Db	301	CCTGATCGCGCTCTGGTGTGGCGACCAAGSGCTGGAAGCGAAACCGGACGTCTGTTA	360
QY	361	CAGGACGTGGCGCTGAGGCCTTAGCGGATCAAAATTCGCTGGCGGTTATCTTGCCCA	420
Db	361	CAGGACGTGGCGCTGAGGCCTTAGCGGATCAAAATTCGCTGGCGGTTATCTTGCCCA	420
QY	421	ACGTTTGCAAGAACTGGCGGCAGGTTTACGACAGCTATTTGCTGGCTCGACCGAT	480
Db	421	ACGTTTGCAAGAACTGGCGGCAGGTTTACGACAGCTATTTGCTGGCTCGACCGAT	480
QY	481	CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGCGGCAAAAGTTCCGCGTTTAC	540
Db	481	CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGCGGCAAAAGTTCCGCGTTTAC	540
QY	541	AGCAATCCGGATTTCATTGGCGTGCAGCTTGCGGCGCGGTTGAAACCGTTATTGCCATT	600
Db	541	AGCAATCCGGATTTCATTGGCGTGCAGCTTGCGGCGCGGTTGAAACCGTTATTGCCATT	600
QY	601	GGTGGGGGATGTCGACCGGTATCGGTTTTGGTGCGAATGCGCGTACGGCGTGATCACC	660
Db	601	GGTGGGGGATGTCGACCGGTATCGGTTTTGGTGCGAATGCGCGTACGGCGTGATCACC	660
QY	661	CGTGGGCTGGCTGAATGTCCGCTCTTGGTGGCGCGCTGGTGCCGACCTGCCACTTT	720
Db	661	CGTGGGCTGGCTGAATGTCCGCTCTTGGTGGCGCGCTGGTGCCGACCTGCCACTTT	720
QY	721	ATGGGCATGGCGGGCTTGGCGATCTGGTGCTTACCTGTACCGAAACAGTCGCGTAAC	780
Db	721	ATGGGCATGGCGGGCTTGGCGATCTGGTGCTTACCTGTACCGAAACAGTCGCGTAAC	780

QY	781	CGCCGTTTTGGCATGATGCTCGGTACGGCATGGATGTACAAAGCCGCGAGAGAAGATT	840
Db	781	CGCCGTTTTGGCATGATGCTCGGTACGGCATGGATGTACAAAGCCGCGAGAGAAGATT	840
QY	841	GGTCAGGTGGTGAAGGCTACCGCAATACGAAAGAAAGTCCGGAACCTGGCGCATCGCTTC	900
Db	841	GGTCAGGTGGTGAAGGCTACCGCAATACGAAAGAAAGTCCGGAACCTGGCGCATCGCTTC	900
QY	901	GGCGTTGAAATGCAATAACCCGAGGAAATTTATCAAGTATTATTTGGGAAAAAACGCG	960
Db	901	GGCGTTGAAATGCAATAACCCGAGGAAATTTATCAAGTATTATTTGGGAAAAAACGCG	960
QY	961	CGCGAGGCGCAGATTGACTTTACTAGTCTGTCACGCAAGGACGAGCGCAGCCACTAA	1020
Db	961	CGCGAGGCGCAGATTGACTTTACTAGTCTGTCACGCAAGGACGAGCGCAGCCACTAA	1020

RESULT 4

ADT48853

ID ADT48853 standard; cDNA; 1020 BP.

XX AC ADT48853;

XX DT 02-DEC-2004 (first entry)

XX DE Bacterial polynucleotide #23604.

XX KW Recombinant DNA construct; transformed plant; improved plant property;
XX cold tolerance; heat tolerance; drought tolerance; herbicide; osmosis;
KW pathogen tolerance; pest tolerance; plant disease resistance;
KW cell cycle pathway modification; plant growth regulator;
KW homologous recombination; seed oil yield; protein yield; carbohydrate;
KW nitrogen; phosphorus; photosynthesis; lignin; galactomannan;
KW bacterial polynucleotide; gene; ss.

OS Bacteria.

XX US2003233675-A1.

XX PD 18-DEC-2003.

XX PF 20-FEB-2003; 2003US-00369493.

XX PR 21-FEB-2002; 2002US-0360039P.

XX PA (CAOY/) CAO Y.
PA (HINK/) HINKLE G J.
PA (SLAT/) SLATER S C.
PA (CHEN/) CHEN X.
PA (GOLD/) GOLDMAN B S.

PI Cao Y, Hinkle GJ, Slater SC, Chen X, Goldman BS;

XX WPI; 2004-061375/06.

PT New recombinant DNA construct comprising a promoter positioned to provide
PT for expression of a polynucleotide encoding a polypeptide from a
PT microbial source, useful for producing plants with improved properties.

PS Claim 1; SEQ ID NO 47291; 122pp; English.

XX The invention relates to a recombinant DNA construct comprising a
CC promoter functional in a plant cell, where the promoter is positioned to
CC provide for expression of a polynucleotide encoding a polypeptide from a
CC microbial source. The invention also relates to a transformed plant
CC comprising the recombinant DNA construct and a method of producing a
CC transformed plant having an improved property. The plant is a crop plant
CC such as maize or soybean. The method of producing a transformed plant
CC having an improved property comprises transforming a plant with the
CC recombinant DNA construct and growing the transformed plant, where the
CC polynucleotide or polypeptide is useful for improving plant properties.
CC The recombinant DNA construct is useful for producing plants with
CC improved plant properties, e.g. improved cold, heat or drought tolerance,

CC tolerance to herbicides, extreme osmotic conditions, pathogens or pests,
CC increased resistance to plant disease, better growth rate by modification
CC of the cell cycle pathway with plant growth regulators, increased rate of
CC homologous recombination, modified seed oil or protein yield and/or
CC content, improved yield by modification of carbohydrate, nitrogen or
CC phosphorus use and/or uptake, by modification of photosynthesis or by
CC providing improved plant growth and development under at least one stress
CC condition, improved lignin production or improved galactomannan
CC production. This sequence represents a bacterial polynucleotide used in
CC the scope of the invention. Note: The sequence data for this patent did
CC not form part of the printed specification but was obtained in electronic
CC format from USPTO at seqdata.uspto.gov/sequence.html.
XX
SQ Sequence 1020 BP; 213 A; 275 C; 304 G; 228 T; 0 U; 0 Other;

Query Match 99.8%; Score 1018.4; DB 13; Length 1020;
Best Local Similarity 99.9%; Pred. No. 6.8e-308;
Matches 1019; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCGGCTCGTACGGCACCGCTCTT 60
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCGGCTCGTACGGCACCGCTCTT 60
QY 61 GCCATCACCTGGCAAGAAATGGCCACGAGTTGCTCTCTGGGGCCATGACCCCTGAACAT 120
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
61 GCCATCACCTGGCAAGAAATGGCCACGAGTTGCTCTCTGGGGCCATGACCCCTGAACAT 120
QY 121 ATCGCAACGCTTGAAACGCGACCGCTGTAACGCCCGCTTCTCCCGATGTGCCCTTTCCC 180
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
121 ATCGCAACGCTTGAAACGCGACCGCTGTAACGCCCGCTTCTCCCGATGTGCCCTTTCCC 180
QY 181 GATACGCTCCATCTTGAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAATATCTC 240
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
181 GATACGCTCCATCTTGAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAATATCTC 240
QY 241 GTCGTGTAACCCAGCCATGCTTTGGTGAAGTGCTGCGCAGATTAAACCACTGATCGGT 300
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
241 GTCGTGTAACCCAGCCATGCTTTGGTGAAGTGCTGCGCAGATTAAACCACTGATCGGT 300
QY 301 CCTGATCGCGCTCTGGTGTGGCGCACCAAGGGCTGGAAGCGGAAACCGGACGCTGTGTTA 360
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
301 CCTGATCGCGCTCTGGTGTGGCGCACCAAGGGCTGGAAGCGGAAACCGGACGCTGTGTTA 360
QY 361 CAGGACGTGGCGCTGAGGCTTAGCGCATCAAAATTCGCTGGCGGTTATCTTGGCCCCA 420
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
361 CAGGACGTGGCGCTGAGGCTTAGCGCATCAAAATTCGCTGGCGGTTATCTTGGCCCCA 420
QY 421 ACGTTTGCAGAAAGAACTGGCGGCAGGTTTACCGACAGCTATTTGCTGGCCTCGACCGAT 480
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
421 ACGTTTGCAGAAAGAACTGGCGGCAGGTTTACCGACAGCTATTTGCTGGCCTCGACCGAT 480
QY 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGGGCAAAAGTTTCCGCGTTTAC 540
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGGGCAAAAGTTTCCGCGTTTAC 540
QY 541 AGCAATCCGGATTTCAATTGGCGTGCAGCTTGGCGGCGCGGTGAAAAACGTTATTGCCATT 600
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
541 AGCAATCCGGATTTCAATTGGCGTGCAGCTTGGCGGCGCGGTGAAAAACGTTATTGCCATT 600
QY 601 GGTGCGGGGATGTCCGACGGTATCGGTTTGGTGCAGATCGCGGTACGGCGCTGATCACC 660
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
601 GGTGCGGGGATGTCCGACGGTATCGGTTTGGTGCAGATCGCGGTACGGCGCTGATCACC 660
QY 661 CGTGGGCTGGCTGAAATGTCCGCTCTTGGTGCAGGCGGTGGGTGCCACCTTT 720
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
661 CGTGGGCTGGCTGAAATGTCCGCTCTTGGTGCAGGCGGTGGGTGCCACCTTT 720
QY 721 ATGGGCATGGCGGGCTTGGCGATCTGGTCTTACCTGTACCGGAAACCAAGTCGCGTAAC 780
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
721 ATGGGCATGGCGGGCTTGGCGATCTGGTCTTACCTGTACCGGAAACCAAGTCGCGTAAC 780
QY 781 CGCCGTTTTGGCATGATCTCGGTGAGGCGATGGATGTACAAAGCGCGCAGGAGAGATT 840
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
781 CGCCGTTTTGGCATGATCTCGGTGAGGCGATGGATGTACAAAGCGCGCAGGAGAGATT 840

Dd 781 CGCCGTTTTGGCATGATGCTCGGTGAGGCGATGGATGTACAAAGCGCGCAGGAGAGATT 840
QY 841 GGTGAGGTGGTGAAGGCTACCGCAATACGAAAGAAAGTCCGGAAGTGGCGCATCGCTTC 900
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
841 GGTGAGGTGGTGAAGGCTACCGCAATACGAAAGAAAGTCCGGAAGTGGCGCATCGCTTC 900
QY 901 GGCCTTGAATGCCAATAACCGAGGAAATTTATCAAGTATTATTTGCGAAAAACGCG 960
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
901 GGCCTTGAATGCCAATAACCGAGGAAATTTATCAAGTATTATTTGCGAAAAACGCG 960
QY 961 CGCGAGGCAGCATTTGACTTTTACTAGGTCGTCACGCAAGGACGAGCGCAGCCACTAA 1020
Dd ||||||||||||||||||||||||||||||||||||||||||||||||||||||||||||
961 CGCGAGGCAGCATTTGACTTTTACTAGGTCGTCACGCAAGGACGAGCGCAGCCACTAA 1020

RESULT 5

ACA51335

ID ACA51335 standard; DNA; 1020 BP.

XX ACA51335;

DT 19-JUN-2003 (first entry)

XX Prokaryotic essential gene #32992.

DE Antisense; ds; prokaryotic essential gene; cell proliferation;
XX drug design; gene.

OS Salmomella typhi.

XX WO200277183-A2.

XX 03-OCT-2002.

PF 21-MAR-2002; 2002WO-US009107.

XX 21-MAR-2001; 2001US-00815242.

PR 06-SEP-2001; 2001US-00948993.

PR 25-OCT-2001; 2001US-0342923P.

PR 08-FEB-2002; 2002US-00072851.

PR 06-MAR-2002; 2002US-0362699P.

XX (ELIT-) ELITRA PHARM INC.

XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;

XX WPI; 2003-029926/02.

DR P-PSDB; ABU47465.

XX New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.

PS Claim 14; SEQ ID NO 39205; 1766pp; English.

XX The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)
CC identifying a gene required for cellular proliferation or the biological
CC pathway in which a proliferation-required gene or its gene product lies
CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a

CC or a gene on which the test compound that inhibits proliferation of an
CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
CC compound's activity; (11) a culture comprising strains in which the gene
CC product is overexpressed or underexpressed; (12) determining the extent
CC to which each of the strains is present in a culture or collection of
CC strains; or (13) identifying the target of a compound that inhibits the
CC proliferation of an organism. The antisense nucleic acids are useful for
CC identifying proteins or screening for homologous nucleic acids required
CC for cellular proliferation to isolate candidate molecules for rational
CC drug discovery programs, or for screening homologous nucleic acids
CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is one of the target
CC prokaryotic essential genes. Note: The sequence data for this patent did
CC not form part of the printed specification, but was obtained in
CC electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences
XX

SQ Sequence 1017 BP; 207 A; 289 C; 312 G; 209 T; 0 U; 0 Other;

Query Match 75.1%; Score 765.8; DB 8; Length 1017;
Best Local Similarity 84.6%; Pred. No. 8.6e-229;
Matches 860; Conservative 0; Mismatches 157; Indels 0; Gaps 0;

QY 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCGGCTCGTACGGCACCGCTCTT 60
Db |||||
QY 61 GCCATCACCTGGCAAGAAATGGCCACGAGTTGTCTCTGGGGCCATGACCCGTGAACAT 120
Db |||||
QY 121 ATCGCAACGCTTGAAACGCGACCGCTGTAAACGCCGCGTTTCTCCCGATGTCCTTTCCC 180
Db |||||
QY 181 GATACGCTCCATCTTGAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAAATATTCTC 240
Db |||||
QY 241 GTCGTCGTACCCAGCCATGCTTTGGTGAAGTGTCTGGCCAGATTAAACCACTGATCGGT 300
Db |||||
QY 301 CCTGATCGCGCTCTGGTGTCGGCGCACCAAGGGCTGGAAGCGGAAACCGGACGCTCTGTTA 360
Db |||||
QY 361 CAGGACGTGGCGGTGAGGCTTAGGCGATCAAAATCCGCTGGCGGTATCTCTGGCCCCA 420
Db |||||
QY 421 ACGTTTGCAGAAAGAACTGGCGGCAGGTTTACCGACAGTATTTTCGCTGGCCTCGACCGAT 480
Db |||||
QY 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACCTGCGGCAAAAGTTTCCGCGTTTAC 540
Db |||||
QY 541 AGCAATCCGGATTTTCATTTGGCGTGCAGCTTGGCGGCGCGGTGAAAAACGTTATTGCCATT 600
Db |||||
QY 601 GGTGCGGGGATGTCGACGCGTATCGGTTTGGTGCGAATGCGCGTACGGCGCTGATCACC 660
Db |||||
QY 661 CGTGGGCTGGCTGAAATGTCGCGTCTTGGTGCGGCGCTGGGTGCCAGCCCTGCCACCTTT 720
Db |||||
QY 721 ATGGCATGGCGGGCTTGGCGATCTGGTGCTTACCTGTACCGGAAACCGAGTCGGGTAAC 780
Db |||||

Db 721 ATGGGAATGGCTGGCTGGGCGACCTGGTGTGCTGACCTGTACCGATAACCACTCTCGTAAC 780
QY 781 CGCCGTTTTGGCATGATGCTCGGTGAGGCGATGATGTACAAAGCGCGCAGGAGAAGATT 840
Db |||||
QY 841 GGTGAGGTGGTGAAGGCTACCGCAATACGAAAGAGTCCGCGAATGTCGCGCATCGCTTC 900
Db |||||
QY 901 GCGGTTGAAATGCCAATAACCGAGGAAATTTATCAAGTATTATATTCGGGAAAAACGCG 960
Db |||||
QY 961 GCGGAGGCAGCATTTGACTTTACTAGTCTGTCACGCAAGGACGAGCGCAGGCCAC 1017
Db |||||

RESULT 7
ACA35838

ID ACA35838 standard; DNA; 1017 BP.

XX ACA35838;

DT 19-JUN-2003 (first entry)

XX Prokaryotic essential gene #17495.

DE Antisense; ds; prokaryotic essential gene; cell proliferation;
XX drug design; gene.

OS *Klebsiella pneumoniae*.

XX WO200277183-A2.

PN 03-OCT-2002.

PF 21-MAR-2002; 2002WO-US009107.

XX 21-MAR-2001; 2001US-00815242.

PR 06-SEP-2001; 2001US-00948993.

PR 25-OCT-2001; 2001US-0342923P.

PR 08-FEB-2002; 2002US-00072851.

XX 06-MAR-2002; 2002US-0362699P.

PA (ELIT-) ELITRA PHARM INC.

XX Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;

XX WPI; 2003-029926/02.
DR P-PSDB; ABU31968.
XX New antisense nucleic acids, useful for identifying proteins or screening
PT for homologous nucleic acids required for cellular proliferation to
PT isolate candidate molecules for rational drug discovery programs.

PS Claim 14; SEQ ID NO 23708; 1766pp; English.

XX The invention relates to an isolated nucleic acid comprising any one of
CC the 6213 antisense sequences given in the specification where expression
CC of the nucleic acid inhibits proliferation of a cell. Also included are:
CC (1) a vector comprising a promoter operably linked to the nucleic acid
CC encoding a polypeptide whose expression is inhibited by the antisense
CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
CC polypeptide or its fragment whose expression is inhibited by the
CC antisense nucleic acid; (4) an antibody capable of specifically binding
CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
CC proliferation or the activity of a gene in an operon required for
CC proliferation; (7) identifying a compound that influences the activity of
CC the gene product or that has an activity against a biological pathway
CC required for proliferation, or that inhibits cellular proliferation; (8)

QY 1 ATGAACCAACGTAATGCTTCAATGACTGTGATCGGTGCCGGCTCGTACGGCACCGCTCTT 60
| | | | |
Db 1 ATGAACCAACCAACCTGCTTCAATGGCTGTTATCGGTGCCGATCTTACGGCACCGCATTA 60
| | | | |
QY 61 GCCATCACCTGGCAAGAAATGGCCACGAGGTTGCTCTGTGGGCCATGACCCCTGAACAT 120
| | | | |
Db 61 GCTATCACACTGGCGCGTAATGGCCATCAAGTCGTGTTATGGGCCATGACCCCTAAACAT 120
| | | | |
QY 121 ATCGCAACGCTTGAAACGCGCTGTAACGCGCTGTAACGCGCTTCTCCCCGATGTGCCCTTCCC 180
| | | | |
Db 121 ATTCAACAGCTGCAACAAGACCGCTGTAACCGCGCTTCTACCTGATGCTGCTTTCCCC 180
| | | | |
QY 181 GATACGCTCCATCTTGAAGCGATCTGCCACATCGCTGCGCAGCCAGCCGTAATATCTC 240
| | | | |
Db 181 GATACGTTGCGATTGGAACCGGACTTAGCATGTGCGTTGGTGTGCCCGCGATGTGTG 240
| | | | |
QY 241 GTCGTCGTACCCAGCCATGTCTTTGGTGAAGTGTGCGCCAGATTAACCACTGATGCGT 300
| | | | |
Db 241 GTCGTCGTGCCAGCCATGTCTTTGGTGCTGTTTACATCAGTTGAAGCCTCATCTACGT 300
| | | | |
QY 301 CCTGATCGCGCTCTGCTGTGGCGCACCAAGGGCTGGAAGCGGAACCGGACGTCGTGTTA 360
| | | | |
Db 301 AAAGATGCACGTATCGTCTGGCAACCAAGGGCTAGAGCTGAACCGGCCGCTGCTA 360
| | | | |
QY 361 CAGGACGTGGCGCTGAGGCTTAGCGGATCAAAATTCGCTGGCGGTTATCTCGGCCCA 420
| | | | |
Db 361 CAGGATGTGGCCCGGAAGTCTTGGGCGAGGCTATCCCGCTTGCCGTGATTTCTGCTCCA 420
| | | | |
QY 421 ACGTTTGCAGAAAGAACTGGCGCAGGTTTACCGACAGCTATTTGCTGGCCTCGACCGAT 480
| | | | |
Db 421 ACGTTTGCCAAAGAAATTGGCGCGGGTTTGCCCTACGGCGATTGCGTTGGCATCGACCGAT 480
| | | | |
QY 481 CAGACCTTTGCCGATGATCTCCAGCAGCTGCTGCACTGCGGCAAAAGTTTCCGCGTTTAC 540
| | | | |
Db 481 GTGCAATTAGCGAAGATCTGCAACAGTTATTGCACTGTGGAAGAAAGCTTTTCGAGTTTAC 540
| | | | |
QY 541 AGCAATCCGATTTTCATTTGGCGTGACGCTTGGCGCGCGGCTGAAACGTTATTGCCATT 600
| | | | |
Db 541 AGTAATCCTGATTTTATCGGGGTACAGCTTGGTGGCGCAGTGAAACGTTGATTGCCATC 600
| | | | |
QY 601 GGTGCGGGGATGTCCGACGGTATCGGTTTGGTGCGAATGCGCGTACGGCGGTGATCACC 660
| | | | |
Db 601 GGTGCAGGTATGTCCGATGGCATCGGTTTGGTGCGAATGCCCCGTACCGCTCTAATAACC 660
| | | | |
QY 661 CGTGGGCTGGCTGAAATGTCGCTCTTGGTGCGGCGTGGGTGCCACCTGCCACTTT 720
| | | | |
Db 661 CGCGGTTAGCGGAGATGACGCGCTTAGGGACGGCATTAGGTGCCGATCCTTCCACCTT 720
| | | | |
QY 721 ATGGGCATGGCGGGCTTGGCGATCTGGTGCTTACCTGTAACCGGAAACCAAGTCGCGTAAC 780
| | | | |
Db 721 ATGGGCATGGCAGGTTTAGGCGATTGGTGCTAACCTGCACAGATAACCAATCCCGTAAC 780
| | | | |
QY 781 CGCCGTTTGGCATGATGCTCGGTGAGGCGATGGATGTACAAAGCGCGCAGGAGAAGATT 840
| | | | |
Db 781 CGCCGATTGGCATATTGCTGGTCAAGGTTTGGGGGTGAAGGAGCGCGCAGGACAACATT 840
| | | | |
QY 841 GGTCAAGTGGTGAAGGCTACCGCAATACGAAAGAAAGTCCGCGAACTGGCGCATCGCTTC 900
| | | | |
Db 841 GGTCAAGTGGTAGAAGTTTACCGTAATACCAAGGAAGTTCTGGCATTAGCACAGCGTCAT 900
| | | | |
QY 901 GCGGTTGAAATGCCAATAACGAGGAAATTTATCAAGTATTATATTGCGGAAACGCG 960
| | | | |
Db 901 GCGGTCGAAATGCCAATAACTGAACAAATTTATCAAGTGTGTATTGTATCAAGAATGCT 960
| | | | |
QY 961 CGGAGGCAGCATTGACTTTTACTAGGTGCTGTCACGCAAGGACGAGCGCAGCAGC 1014
| | | | |
Db 961 CGTGAGCGGCTCTGACGTTGTTGGGGCGGACCAAAAAAGATGAAAAAATCGGC 1014
| | | | |

RESULT 11
ABQ21987/c
ID ABQ21987 standard; DNA; 781 BP.

XX ABQ21987;
AC 12-JUL-2002 (first entry)
XX
DT
XX
DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 8578.
XX
KW Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW drug; side effect; cancer; central nervous system; cardiovascular;
KW gastrointestinal; respiratory system; single nucleotide polymorphism;
KW SNP; cell differentiation; ds.
XX
OS Homo sapiens.
XX
PN WO200218632-A2.
XX
PD 07-MAR-2002.
XX
PF 01-SEP-2001; 2001WO-EP010074.
XX
PR 01-SEP-2000; 2000DE-01043826.
PR 05-SEP-2000; 2000DE-01044543.
XX
PA (EPIG-) EPIGENOMICS AG.
XX
PI Olek A, Piepenbrock C, Berlin K, Guetig D;
XX
DR WPI; 2002-371829/40.
XX
PT Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons
PT from chemically treated DNA.
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
CC This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one member,
CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
CC degree of hybridisation to both classes is determined from the label on
CC the amplicon. From the ratio of labels hybridised to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:
CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's); and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation
CC status of many C residues to be determined simultaneously. ABQ13410-
CC ABQ54121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the
CC disclosure of the invention
XX
SQ Sequence 781 BP; 313 A; 239 C; 76 G; 153 T; 0 U; 0 Other;
Query Match 55.5%; Score 565.6; DB 6; Length 781;
Best Local Similarity 82.8%; Pred. No. 3.7e-166;
Matches 646; Conservative 0; Mismatches 134; Indels 0; Gaps 0;
QY 153 CGCGTTTCTCCCCGATGTCCTTTTCCCGATACGCTCCATCTTGAAAGCGATCTCGCCAC 212
| | | | |
Db 780 CGCGTTTTTTTCGATGTGTTTTTATTTCGATACGTTTATTTTGAAAGTAATTTTCGTAT 721
| | | | |
QY 213 TCGCTGGCAGCCAGCCGTAATATTCTCGTCGTCGTACCCAGCCATGTCTTTTGGTGAAGT 272
| | | | |
Db 720 TCGGTTGGTAGTTAGTCGTAATATTTTCGTCGTCGTTATTTAGTTATGTTTTTGGTGAAGT 661
| | | | |
QY 273 GCTGCGCCAGATTAAACCACTGATGCGTCTCTGATGCGCGTCTGGTGTGGCGCAAAAGG 332
| | | | |
Db 660 GTTGCGTTAGATTAAATTATTGATGCGTTTTTGATGCGCGTTTGGTGTGGCGGATTAAAGG 601
| | | | |

WP	ACF67367_03	300001	410000	
WP	ACF67367_04	400001	510000	
WP	ACF67367_05	500001	610000	
WP	ACF67367_06	600001	710000	
WP	ACF67367_07	700001	810000	
WP	ACF67367_08	800001	910000	
WP	ACF67367_09	900001	1010000	
WP	ACF67367_10	1000001	1110000	
WP	ACF67367_11	1100001	1210000	
WP	ACF67367_12	1200001	1310000	
WP	ACF67367_13	1300001	1410000	
WP	ACF67367_14	1400001	1510000	
WP	ACF67367_15	1500001	1610000	
WP	ACF67367_16	1600001	1710000	
WP	ACF67367_17	1700001	1810000	
WP	ACF67367_18	1800001	1910000	
WP	ACF67367_19	1900001	2010000	
WP	ACF67367_20	2000001	2110000	
WP	ACF67367_21	2100001	2210000	
WP	ACF67367_22	2200001	2310000	
WP	ACF67367_23	2300001	2410000	
WP	ACF67367_24	2400001	2510000	
WP	ACF67367_25	2500001	2610000	
WP	ACF67367_26	2600001	2710000	
WP	ACF67367_27	2700001	2810000	
WP	ACF67367_28	2800001	2910000	
WP	ACF67367_29	2900001	3010000	
WP	ACF67367_30	3000001	3110000	
WP	ACF67367_31	3100001	3210000	
WP	ACF67367_32	3200001	3310000	
WP	ACF67367_33	3300001	3410000	
WP	ACF67367_34	3400001	3510000	
WP	ACF67367_35	3500001	3610000	
WP	ACF67367_36	3600001	3710000	
WP	ACF67367_37	3700001	3810000	
WP	ACF67367_38	3800001	3910000	
WP	ACF67367_39	3900001	4010000	
WP	ACF67367_40	4000001	4110000	
WP	ACF67367_41	4100001	4210000	
WP	ACF67367_42	4200001	4310000	
WP	ACF67367_43	4300001	4410000	
WP	ACF67367_44	4400001	4510000	
WP	ACF67367_45	4500001	4610000	
WP	ACF67367_46	4600001	4710000	
WP	ACF67367_47	4700001	4810000	
WP	ACF67367_48	4800001	4910000	
WP	ACF67367_49	4900001	5010000	
WP	ACF67367_50	5000001	5110000	
WP	ACF67367_51	5100001	5210000	
WP	ACF67367_52	5200001	5310000	
WP	ACF67367_53	5300001	5410000	
WP	ACF67367_54	5400001	5510000	
WP	ACF67367_55	5500001	5610000	
WP	ACF67367_56	5600001	5648894	

Query Match

Best Local Similarity

Matches 722; Conservative

54.5%; 72.3%; 0;

Score 556.4; DB 10; Length 110000;

Pred. No. 3.1e-162;

Mismatches 276; Indels 0; Gaps 0;

QY	11	GTAATGCTTCAATGACTGTGATCGGTGCCGGCTCGTACGGCACCGCTCTTGGCATCACCC	70
Db	79833	GTACTGTTTCTATGACAGTGAATGGTGTGCTCATACGGCACCTCATTAGCCATTACGC	79892

QY	71	TGGCAAGAAATGGCCACGAGTGTGCTCTGCGGCCATGACCCCTGAACATATCGAACGC	130
Db	79893	TGGCTCGTAATGGTTCATAATGTTGTACTTTGGGGGCATAATCCAGAGCATGTTGGGGCAT	79952

QY	131	TTGAACGCGACCGCTGTACGCGCGGTTTCTCCCGATGCGCTTTTCCCGATACGCTCC	190
Db	79953	TGCAACGGGTGCGTGTGTAATCAAAAAATTCTGCCGGATGTTCCCTTCCCTGATAGTTAT	80012

QY	191	ATCTTGAAAGCGATCTCGCCACTGCGCTGGCAGCCAGCCGTAATATCTCGTCGCTAC	250

Db	80013	TGCTTGAAACGGACCTAATAAAAAAGCACATAACAGCGAGCCGCGATATTCTTGTGTGTGTAC	80072
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QY	251	CCAGCCATGTCTTTGGTGAAGTGTCTGCGCCAGATTAAACCACATGATGCGTCTGATGCGC	310
Db	80073	CTAGCCATGTGTTTGGTGAAGTGTAAAGCAGATAAAAAACCACATTTACGGCCTGATTAC	80132

QY	311	GTCTGGTGTGGCGACCAAAAGGCTGGAAGCGGAAACCGGACGCTCTGTTACAGGACGTGG	370
Db	80133	GTATCGTATGGGCAACTAAAGGCTTGAAGCGGATACCGTGGTTATTGACAGGATGTGG	80192

QY	371	CGCGTGAGGCCCTTAGCGCATCAAATTCGCTGGCGGTTATCTCTGGCCCCAACGTTTCCGA	430
Db	80193	CCCGTGAGATATTAGGCAATGAAATACCGCTAGCGGTGCTCTCTGGGGCCAAACATTTGCTA	80252

QY	431	AAGAACTGGCGGCGAGGTTTACCGACAGCTATTTTCGCTGGCCTCGACCGATCAGACCTTTG	490
Db	80253	AAGAGTTAGCGGCTGGTTTGCCCTACCGCGATTGCTATTTCCGCGACGGAATCTGCTTTTG	80312

QY	491	CCGATGATCTCCAGCAGCTGCTGCACTGCGGCGCGGTGAAAAACGTTATTGCCATTGGTGGGGA	550
Db	80313	GCGATGGACTTCAACAATATTCCACTGTGGCAAAAGTTTCCGGGTTTATAAAAAATCCTG	80372

QY	551	ATTTCATTGGCGTGCACTTGGCGCGCGGTGAAAAACGTTATTGCCATTGGTGGGGA	610
Db	80373	ATTTTATTGGTGTTCAACTCGGTGGTGCCGTAAAAAACGTCATCGCCATTGGCGCGGGA	80432

QY	611	TGTCCGACGGTATCGGTTTGGTGCGAATGCGCGTACGGCGCTGATCACCCGCTGGGCTGG	670
Db	80433	TATCTGATGGCATGGGATTTGGTGCTAAATGCTCGTACCGCATTGATTACTCGTGGATTGG	80492

QY	671	CTGAAATGTCGCGTCTTGGTGCGGCGCTGGGTGCCGACCCCTGCCACCTTTTATGGGCATGG	730
Db	80493	CGGAAATGAGTCGCCCTTGGTGACGCGCTTGGTGTGATCCTTCTACCTTTATGGGCATGG	80552

QY	731	CGGGGCTTGGCGATCTGGTGCTTACCTGTACCGAAAAACCAGTCGCGTAACCGCCGTTTGG	790
Db	80553	CGGGATTGGGCGATTGGTCTTAACTTGTACTGATAAACCAATCACGTAACCGTCGTTTG	80612

QY	791	GCATGATGCTCGGTACGGGCATGGATGTACAAAGCGCGCAGGAGAAGATTGGTCAGGTGG	850
Db	80613	GCATGATGCTGGGCGAGGGAATCAGTGTGGAAGAAGCGCAGTATCAGATTGGGCGAGTTG	80672

QY	851	TGGAAGGCTACCGCAATACGAAAGAAGTCCGCGAACTGGCGCATCGCTTCGCGCTTGAAA	910
Db	80673	TTGAAGGTTATCGCAATACCAAGAAGTACGTGCAATTGGCTAATCGCGCCAAATGTAGAAA	80732

QY	911	TGCCAATAACCGAGGAAATTTATCAAGTATTATATTTCGGGAAAAAACCGCGCGAGGCAG	970
Db	80733	TGCCGATTGCAGAACAAATCTACCAGATATCTCTATTGCAATAAAAAATGTATAGAGCTG	80792

QY	971	CATTGACTTTTACTAGTGTGTCACGCAAGGACGAGCGC	1008
Db	80793	CTCAGGCATTATTAGGAAGAGCCAGAAAGGATGAGAGC	80830

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Job time : 666 secs

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